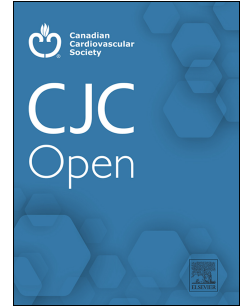


Journal Pre-proof

Adherence to cardiovascular prevention guidelines in an academic center

Iness Soltani, MD, Marie-Claude Beaulieu, MD, Maude Sestier, MD, Hao Cheng Shen, MD, Ali Hillani, MD, Alexis Matteau, MD SM, Samer Mansour, MD, Brian J. Potter, MDCM SM



PII: S2589-790X(23)00064-1

DOI: <https://doi.org/10.1016/j.cjco.2023.03.010>

Reference: CJCO 653

To appear in: *CJC Open*

Received Date: 25 February 2023

Revised Date: 18 March 2023

Accepted Date: 20 March 2023

Please cite this article as: I. Soltani, M.-C. Beaulieu, M. Sestier, H.C. Shen, A. Hillani, A. Matteau, S. Mansour, B.J. Potter, Adherence to cardiovascular prevention guidelines in an academic center, *CJC Open* (2023), doi: <https://doi.org/10.1016/j.cjco.2023.03.010>.

This is a PDF file of an article that has undergone enhancements after acceptance, such as the addition of a cover page and metadata, and formatting for readability, but it is not yet the definitive version of record. This version will undergo additional copyediting, typesetting and review before it is published in its final form, but we are providing this version to give early visibility of the article. Please note that, during the production process, errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

© 2023 Published by Elsevier Inc. on behalf of the Canadian Cardiovascular Society.

1 **Adherence to cardiovascular prevention guidelines in an academic center**

2

3

4

5

6 Iness Soltani MD^{#,1} Marie-Claude Beaulieu MD^{#,1}, Maude Sestier MD¹,

7 Hao Cheng Shen MD¹, Ali Hillani MD¹, Alexis Matteau MD SM¹, Samer Mansour MD¹, &

8 Brian J. Potter MDCM SM¹

9

10 **#. Co-First authors with equal contribution.**

11 1. Centre hospitalier de l'Université de Montréal (CHUM) Research Center and
12 Cardiovascular Center, Montreal, Canada

13

14 Short title: Cardiovascular prevention in an academic center

15 Brief Summary: 60 words

16 Abstract: 250 words

17 Text: 3233 words

18 Tables: 2

19 Supplemental tables: 7

20

21

22 Author of correspondence:

23

24 Brian J. Potter, MDCM SM FRCPC

25 Carrefour de l'innovation et évaluation en santé (CIÉS),

26 Centre de recherche du CHUM (CRCHUM)

27 Cardiology & Interventional Cardiology, CHUM

28 Pavillon S, S03-334

29 850, rue St-Denis

30 Montréal, QC, Canada

31 H2X 0A9

32 brian.potter@umontreal.ca

33 Tel: 514-890-8000 ext.15473

34 Fax: 514-412-7212

35 Brief summary

36

37 Cardiovascular prevention is essential to reduce the morbidity and mortality of cardiovascular
38 diseases. Our study aims to identify the practice gaps in the adherence to cardiovascular
39 prevention guidelines among cardiologists. This retrospective study reveals a suboptimal
40 adherence to cardiovascular prevention guidelines for both pharmacological and non-
41 pharmacological interventions. Strategies such as professional development programs and
42 application-based decision support are potential solutions.

43

Journal Pre-proof

44 Abstract

45

46 Background

47 Adherence to guidelines is associated with better patient outcomes. While studies show
48 suboptimal adherence to cardiovascular prevention guidelines among general practitioners,
49 adherence among specialist physicians is understudied. The aim of this analysis was to identify
50 practice gaps among cardiologists in a tertiary academic center.

51

52 Methods

53 We retrospectively audited cardiology outpatient clinic notes at the *CHUM* from January 1st,
54 2019 to February 28th, 2019. Data were abstracted from hospital medical records. The primary
55 outcome of interest was the rate of adherence to cardiovascular prevention guidelines. We
56 compared the chart documented practice at our center to the Canadian hypertension, lipid,
57 diabetes, antiplatelet and heart failure guidelines in effect at the time of the audit. We also
58 collected information regarding discussions of smoking, alcohol consumption, physical
59 activity and diet.

60

61 Results

62 A total of 2503 patients were included with a mean age of 65.6±14.5 years. Dyslipidemia
63 occurred in 63%, hypertension in 55% and coronary artery disease in 41%. Optimal LDL
64 control was documented achieved in just 39% of cases. Blood pressure control was adequate
65 for 65% of patients, and glycemic control was achieved in 47% of patients with diabetes. Heart
66 failure treatment was optimal in 34%. Nearly all (95%) patients with CAD had appropriate
67 antithrombotic therapy. Discussion of non-pharmacologic interventions varied from 91%
68 (smoking) to 16% (diet).

69

70 Conclusions

71 Primary and secondary prevention of cardiovascular events was found to be suboptimal in an
72 academic tertiary care outpatient cardiology clinic and may be representative of similar
73 shortcomings nationwide. Strategies to ensure guideline adherence are needed.

74

75

76 Introduction

77 Cardiovascular disease (CVD) has been the leading cause of death in North America for many
78 years.^{1,2} Morbidity and mortality associated with CVD represent significant costs and burden
79 of care for the healthcare system globally.³ As it is well known that adverse cardiovascular
80 events can be prevented through healthy behaviors and optimal treatment of cardiovascular risk
81 factors,⁴ national scientific societies have published evidence-based guidelines for the
82 prevention of CVD events through management of diabetes, lipid disorders, hypertension, heart
83 failure, coronary artery disease (CAD) and peripheral artery disease (PAD), including both
84 pharmacologic and non-pharmacologic interventions.⁴⁻¹² The American College of Cardiology
85 (ACC) also recommends a team-based approach to optimize preventive cardiology aspects and
86 minimize CVD.⁴

87 Despite strong evidence that adherence to prevention guidelines is associated with better
88 outcomes for patients,^{4, 13-16} studies have shown that adherence to these guidelines remains
89 suboptimal.^{13, 17-22} For example, adherence rate to individual guideline recommendations was
90 found to range between 5% to 34% even among patients with prior myocardial infarction.²¹
91 The Million Hearts study also assessed quality of adherence to preventive cardiology concepts.
92 Antiplatelet prescription, hypertension control, hyperlipidemia control amongst patients with
93 diabetes and tobacco use screening and intervention were appropriate according to guidelines
94 in 71.9%, 66.6%, 75.8% and 79.8% of cases, respectively.¹⁸

95 Despite the availability of well-developed cardiovascular prevention guidelines, a number
96 of barriers appear to limit their application in clinical practice.²³ However, while guideline
97 adherence has been evaluated among general practitioners,¹⁷⁻²¹ there is limited data regarding
98 guideline adherence among cardiologists.²⁴ Better defining treatment gaps among specialists
99 may shed important light on the nature of these barriers to improve real-world prevention and
100 lead to specific interventions.

101

102 Methods

103 This study consisted of a retrospective chart audit of all patients ≥ 18 years of age seen by a
104 cardiologist in the outpatient Cardiology Clinic at the *Centre hospitalier de l'Université de*
105 *Montréal* (CHUM) from January 1st, 2019, to February 28th, 2019. This period was chosen as
106 it allowed a second follow-up audit of the same period in 2020 prior to the COVID-19
107 pandemic reaching North America. Cardiologist visits occurring both in general cardiology and
108 specialized cardiology clinics were included.

109 The CHUM is a large Canadian tertiary care academic center that uses the OACIS electronic
110 medical record (EMR) system (Telus Santé, Montréal, Québec, Canada). Outpatient cardiology
111 notes can be made using either a structured digital form or a hand-written note (subsequently
112 scanned into the EMR) according to individual physician preference. The primary outcome of
113 interest was adherence to cardiovascular prevention guidelines in an outpatient setting. We
114 compared the documented practice at our center to the most recent Canadian diabetes, lipids,
115 antiplatelets, hypertension, and heart failure guidelines published at the time of the patient
116 visits.⁴⁻¹²

117 Patients who consulted the Cardiology service more than once during the study period were
118 analyzed based on their more recent visit. Visit notes by non-cardiologist health professionals
119 were not audited (the vast majority of patients were seen only by their cardiologist during the
120 period of study). All data were abstracted from the EMR.

121 For hypertension, we compared charted practice to the 2018 Hypertension Canada
122 guidelines.⁸ Treatment was analyzed for each patient first by collecting medications that were
123 initiated (ACE inhibitor (ACEi), angiotensin receptor neprilysin inhibitor (ANRI), beta-
124 blocker (BB), calcium channel blocker (CCB), diuretics, mineralocorticoid antagonist (MRA),
125 thiazides or angiotensin receptor blocker (ARB)). Blood pressure targets were <130/80 mmHg
126 for patients with diabetes and <140/90 mm Hg for others, needing to reach systolic and diastolic
127 values to be considered on target.

128 Regarding dyslipidemia, we used the 2016 Canadian Cardiovascular Society (CCS)
129 dyslipidemia guidelines and the 2018 CCS familial hypercholesterolemia update.^{6,7} We looked
130 at the rate of lipid treatment target achievement (LDL <2 mmol/L or reduction of 50% of LDL
131 level compared to baseline values), adequacy of screening, and the types of lipid-lowering
132 therapy prescribed to patients with indication of treatment.

133 We used the 2018 Canadian Diabetes Association Guidelines⁵ to assess adherence to
134 diabetes recommendations. Outcomes of interest included the rate of diabetes screening among
135 cardiology patients without diabetes older than 40 years of age or risk factors for diabetes, as
136 well as the rate of glycosylated hemoglobin (A1C) evaluation and adequate glycemic control
137 (A1C \leq 7.0%) among patients with diabetes. We also assessed the use of SGLT-2 inhibitors
138 (SGLT2i) and GLP-1 receptor agonists (GLP1RA); antidiabetic agents with known cardiac
139 benefit.

140 The 2018 CCS antiplatelet therapy guidelines, the 2016 American Heart Association
141 (AHA)/ACC guideline on lower extremity PAD and the 2018 AHA/American Stroke
142 Association Guidelines for management of ischemic stroke¹⁰⁻¹² were all used to assess the

143 adequacy of antiplatelet therapy among CAD and PAD patients. Appropriateness of
144 prescription was defined as prescription of aspirin when there was an indication or omission of
145 prescription when not indicated. For example, patients with stable CAD with an indication for
146 anticoagulation, omission of aspirin prescription was judged to be appropriate. We also
147 assessed if gastroprotective therapy was prescribed according to the 2008 American College of
148 Cardiology Foundation, American College of Gastroenterology and AHA expert consensus
149 recommendations.²⁵ Indications for gastroprotective therapy included the need of aspirin
150 therapy and history of ulcer disease, gastro-intestinal bleeding, dual antiplatelet therapy
151 (DAPT) or concomitant anticoagulant therapy. Gastroprotective therapy was also indicated if
152 more than one of these risk factors are met: age greater than 60 years old, corticosteroid use,
153 dyspepsia or gastroesophageal reflux symptoms. The criterion regarding gastrointestinal
154 symptoms was not taken into account because of the anticipated likelihood that such data might
155 be missing from a cardiologist's visit documentation. We also screened the cohort to determine
156 the proportion of patient who would be eligible for dual pathway inhibition (low-dose
157 rivaroxaban plus aspirin) based on the Rivaroxaban with or without Aspirin in stable
158 Cardiovascular Disease (COMPASS trial²⁶) that was already published at the time, but not yet
159 incorporated into guidelines and *Régis de l'assurance maladie du Québec* (RAMQ)
160 reimbursement criteria (co-existence of CAD and PAD).

161 We referred to the 2017 CCS Heart Failure Guidelines to assess treatment of patients with
162 heart failure.⁹ We collected the left ventricular ejection fraction (LVEF) for all patients. We
163 considered patients with and LVEF $\leq 40\%$ as having heart failure with reduced ejection fraction
164 (HFrEF), for whom optimal medical therapy consisted of at least triple therapy with a BB, an
165 MRA and an ACEi/ARB/ARNI. Patients in sinus rhythm with a resting heart rate more than
166 70 beats per minute despite adequate treatment with BB were expected to receive ivabradine.
167 Treatment with SGLT2i was not included in the heart failure guidelines at the time of the chart
168 audit. If LVEF was $<35\%$ after 3 months of optimal medical therapy, patients were expected
169 to be offered an implantable cardioverter-defibrillator (ICD), plus cardiac resynchronization
170 therapy (CRT) if they were in sinus rhythm with a QRS that was more than 130ms (left bundle
171 branch pattern).

172 For all patients, we also collected any documented information regarding their lifestyle,
173 including smoking, alcohol consumption, physical activity and nutrition. For patients drinking
174 more than the recommended amount of alcohol (>14 drinks per week for men, >11 drinks per
175 week for women),²⁷ we took note of whether the cardiologist addressed this aspect during the
176 patient visit. For active smokers, we verified if a smoking cessation therapy had been

177 prescribed or discussed. We also evaluated if there was a discussion about physical activity
178 and nutritional or referral to a nutritionist. We also recorded whether the body mass index
179 (BMI) was documented or available in the EMR.

180 Continuous data are reported as means and standard deviations or medians and interquartile
181 ranges (IQR), as appropriate, and categorical/binary data are reported as counts and percent
182 proportions. The study protocol was consistent with the ethical guidelines of the 1975
183 Declaration of Helsinki and was conducted in accordance with the Strengthening the Reporting
184 of Observational Studies in Epidemiology (STROBE) guidelines²⁸. The CHUM Research
185 Center institutional ethics board approved the study and provided a waiver for informed
186 consent.

187 **Results**

188 We included 2503 patients seen at the CHUM cardiology clinic from January 1st, 2019 to
189 February 28th, 2019. The mean age was 66 ± 15 years, with 94% of patients being over 40 years
190 old. Sixty percent of patients were men. The most prevalent comorbidities were dyslipidemia
191 (63%), followed by hypertension (55%) and CAD (41%). Active smokers represented 8% of
192 our population (Table 1).

193 Management of hypertension

194 More than half (55%) of patients suffered from hypertension. Of them, 311 (64%) also had
195 diabetes. Optimal blood pressure control was achieved for 62% of all hypertensive patients and
196 66% of patients with diabetes (Table 2). The most frequent medications used were BB (62%
197 for all, 67% for patients with diabetes) followed by CCB (43% for all, 43% for patients with
198 diabetes) and ACEi (36% for all, 39% for patients with diabetes). The use of ARNI in the whole
199 cohort was low (2%) but higher (17%) for patients with HFrEF (Supplemental Table S1 and
200 S6).

201 Management of dyslipidemia

202 1567 patients (63%) had dyslipidemia (Table 1). Lipid treatment was at target dose in 608
203 (39%, Table 2) patients with dyslipidemia. Adequate LDL control was reached in 46% of
204 patients with CAD. Thirty-five patients (3%) with CAD had LDL level above target without
205 any lipid-lowering therapy. Assessment of lipid levels was documented at least once in the last
206 5 years in 66% of patients with known CAD, 52% with PAD, 62% with diabetes over the age
207
208
209

210 of 40, and 59% of patients with CKD over the age of 50. PCSK9 inhibitors prescription was
211 low (1%; Supplemental Table S2).

212

213 Management of diabetes

214 Among the entire cohort, 644 patients had diabetes (26%; Table 1). Documentation of HbA_{1c}
215 was found in 37% of patients without diabetes compared to 75% of patients with diabetes
216 within the previous 5 years. Among patients with diabetes, target A1C ($\leq 7.0\%$) was achieved
217 in 47% of cases (Table 2). GLP1RA and SGLT2i were used in 2.0% and 7.0% of patients with
218 diabetes, respectively, and neither were prescribed to patients without diabetes. In CAD or
219 PAD patients with diabetes, SGLT2i were prescribed in 7% and 9% of cases, respectively.
220 GLP1RA were prescribed to 4% of patients with diabetes with BMI ≥ 30 (Supplemental Table
221 S3).

222

223 Management of CAD/PAD

224 Forty-six percent of patients suffered from either CAD and/or PAD. Among the 1032 patients
225 with CAD, 989 (96%) were on antithrombotic or anticoagulant therapy. Of the 821 patients
226 with CAD and an aspirin indication, 782 (95%) had an appropriate prescription. Among the
227 318 patients with PAD, 296 (93%) were on antithrombotic or anticoagulant. Of the 173 patients
228 with PAD and an indication for aspirin, 160 (92%) had an appropriate prescription (Table 2).
229 Ninety-six (4%) patients in our cohort were taking aspirin without a guideline approved
230 indication.

231 A total of 1665 patients were taking an antiplatelet or anticoagulant. In this population,
232 828 (50%) were on single antiplatelet therapy (SAPT), 163 (10%) were on double antiplatelet
233 therapy (DAPT), 584 (35%) were on anticoagulation therapy alone, 80 (5%) were on SAPT
234 plus an anticoagulant and 10 (<1%) were on triple therapy (DAPT plus anticoagulant). Among
235 DAPT patients, 101 (62%) were also prescribed a gastroprotective medication, compared to 36
236 (45%) of patients receiving SAPT plus an anticoagulant. All 10 patients on triple therapy had
237 gastroprotective medication.

238 Among the 1172 patients that suffered from either CAD or PAD in our study, 616 were
239 on ASA alone, who could potentially benefit from the addition of low dose rivaroxaban as per
240 the COMPASS study²⁹ (Supplemental Tables S4, S5).

241

242 Management of heart failure

243 Seven percent of patients had HFrEF (Table 1). Of them, 40% had a NYHA functional class
244 greater than one. Fifty seven percent of patients had an LVEF \leq 35%. Treatment with a BB
245 was frequent (88%). A renin-angiotensin modulator (ACEi or ARB or ARNI) was part of the
246 treatment in 73% of cases. Only 40% of patients were on MRA however. Overall, therefore,
247 only 62 patients (34%) were optimal medical therapy defined as triple therapy (BB + renin-
248 angiotensin modulator + MRA, Table 2). This percentage increased to 49% in patients followed
249 at the specialized heart failure clinic, suggesting that the low overall proportion might have
250 been due to legitimate limiting factors like low blood pressure or marginal renal function. Forty
251 seven percent of patients with LVEF \leq 35% either had and ICD or documentation that ICD had
252 been discussed. CRT devices were implanted in 28% of eligible patients with a large QRS and
253 LVEF \leq 35%. About a quarter (27%) of HFrEF patients were followed in a specialized heart
254 failure clinic (Supplemental Tables S6).

255

256 Non-pharmacological prevention

257 A minority of patients were active smokers (8%, table 1), of whom 91% had a documented
258 discussion or intervention. Of 40 patients with documented excessive alcohol consumption, 36
259 (90%) had their alcohol consumption addressed by their cardiologist. In contrast, physical
260 activity was discussed with 650 patients (26%), and diet was discussed with 223 (16%) patients.
261 Similarly, when considering only obese patients, 27% and 11% received physical activity and
262 nutrition advice, respectively. For patients referred to the specialized preventive cardiology
263 clinic, documentation of physical activity and diet discussions increased to 86% and 57%
264 respectively (Supplemental Table S7).

265

266 **Discussion**

267 Our study evaluated adherence to CVD prevention guidelines among cardiologists in a
268 Canadian tertiary academic center. We demonstrated that there appears to be suboptimal
269 application of many aspects of the guidelines even by academic cardiovascular specialists,
270 based on available EMR.

271 Treatment to target of blood pressure, HbA1C, and LDL levels was disappointing overall.
272 Regarding dyslipidemia, particularly, it is noteworthy that, despite low rates of achieving target
273 LDL in our cohort, add-on molecules to statin therapy such as PCSK-9 inhibitors were rarely
274 prescribed. This may be partially explained by administrative hurdles to obtaining PCSK-9
275 reimbursement in Québec.

276 Rates of appropriate aspirin prescription were overall encouraging. However,
277 prescription of gastroprotection, particularly among patients receiving SAPT plus an
278 anticoagulant requires improvement. Diabetes treatments with known cardiac benefits such as
279 SGLT2i and GLP1RA^{30,31} were rarely prescribed, but also were not included in the guidelines
280 in force at the time of the audited visits. Similarly, studies showing benefits of SGLT2i for
281 heart failure patients³²⁻³⁵ were not published at the time.

282 The results of our study are consistent with other studies regarding CVD prevention
283 among non-specialist practitioners. In the Million Hearts study¹⁸, which included more than
284 100 000 patients in the United States, hypertension control ranged from 49% to 75% compared
285 the rate of 65% in our study. Dyslipidemia control amongst patients with diabetes was in
286 contrast higher than in our population. Though it is not immediately clear why this would be,
287 an hypothesis could be that the CHUM serves a downtown population that frequently doesn't
288 benefit from an identified primary care physician. A large European registry of patients with
289 CAD also similarly showed that less than 60% of patients had good risk factor control in a
290 population care for by both primary care physicians and specialists.¹³ While CHUM patients
291 frequently do not have primary care physicians, patients with diabetes and chronic renal failure
292 patients frequently benefit from concurrent specialist follow-up in those areas.

293 While discussions of smoking cessation and alcohol consumption were frequently
294 documented, discussion of exercise and diet appeared to occur infrequently unless patients
295 were also followed at our specialized prevention cardiology clinic. This could partially be
296 explained by the fact that physicians who refer their patients to a preventive clinic might be
297 more likely to document these discussions. Similarly, HFrEF treatment appeared slightly better
298 among patients followed at our heart failure clinic, but still limited, possibly explained by low
299 blood pressure and poor renal function limiting treatment options. Data regarding contra-
300 indication to or side-effects due to specific treatments was not analyzed because it was too
301 inconsistently documented in the medical chart. While it justify the absence of some treatments
302 for some patients, it would be unlikely to explain the extent of non-adherence that we have
303 observed.

304 Taken together, these observations would seem to support the ACC recommendation for
305 a team-based approach to optimize prevention.⁴ In line with this, an Iranian study found better
306 adherence rates than we observed among academic specialists in an in-patient setting,²⁴ where
307 patients benefit from multidisciplinary care and relatively prolonged or repeated contact with
308 the treating team. Multidisciplinary care may also be a means of compensating for the time
309 pressures of a busy outpatient clinic and allow for more complete assessment of the full

310 spectrum of prevention in cardiology. Despite the potential benefit of lifestyle modifications,³⁶
311 patients were counselled on physical activity and diet in only a minority of patient visits (26%
312 and 16%, respectively) whether the patient was obese or not in our cohort. This might be
313 underestimated as it might have been discussed but not reported in the medical chart. It may
314 also be that physicians do not have the time to adequately address these issues during the visit.
315 A survey of physicians from diverse specialties, including cardiologists, revealed that less than
316 two thirds of physicians were likely to give nutritional or physical activity advice to their
317 patients. However, this was influenced by their perception of a patient's risk.³⁷ In either case,
318 increasing access to multidisciplinary follow-up could lead to substantial improvement in
319 outcomes.

320 Some limitations of the present study should be acknowledged. First, this is a single
321 center analysis which may limit generalizability. However, our results are consistent with
322 similar studies in the literature and extend the findings to specialized cardiology centers. We
323 therefore believe that they are likely indicative of practice in other academic centers. Second,
324 it is possible that guideline adherence is underestimated and that we have instead captured poor
325 documentation of adherence. We unfortunately don't have data allowing comparison between
326 hand-written and digital medical notes. However, given the very low rates of documentation
327 of certain interventions, it appears unlikely that documentation shortcomings explain the
328 entirety of the adherence gaps that we have identified. Third, whether a patient is also followed
329 by a family doctor or another specialist is known to be inconsistently documented in the
330 cardiology clinic chart. It is therefore possible that overall adherence across providers is better
331 than what is reported here. On the other hand, as our population included patients followed in
332 specialized clinics, such as the heart failure clinic, it is possible that the rate of guideline
333 adherence would have been lower if patients exclusively followed in general cardiology clinics
334 would have been lower than what we have observed. Finally, reimbursement criteria for
335 medications vary from province to province, and it is possible that such differences may lead
336 to variations in adherence to certain prevention recommendations.

337 Bearing in mind these limitations, there is clearly an opportunity for quality
338 improvement. In addition to continuing medical education initiatives for both patients and
339 physicians, given typical time pressures and the complexity of patients who consult in
340 academic centers, it is likely that system level interventions are necessary. The most obvious
341 intervention at this level is to broaden the availability and use of multidisciplinary care in the
342 outpatient setting based both on our findings and guideline recommendations.²³ In addition, the
343 randomized BRIDGE cardiovascular prevention study²⁰ showed that a combination of case

344 management, feedback reports and educational materials for physicians and patients lead to
345 increased use of evidence-based therapies. It should also be possible to several current
346 technologies to provide real-time checklists and decision aids to providers while also aiding in
347 the proper documentation of important prevention interventions.

348

349 **Conclusions**

350 Prevention of CVD through lifestyle behavior modification and optimization of cardiovascular
351 risk factors is essential. Our study reveals that despite appropriate guidance and continued
352 medical education efforts, adherence to prevention recommendations remain suboptimal even
353 in an academic cardiology clinic. Combining a multidisciplinary approach with standardized
354 updated algorithms with technology that facilitates both documentation and the patient
355 encounter appear well-positioned to improve patient CVD outcomes.

356

357

358 **Acknowledgments**

359 None.

360

361 **Funding sources**

362 This study was funded in part by unrestricted research grants from Bayer Canada and
363 Boehringer-Ingelheim Canada and an unrestricted educational grant from Novartis Canada.
364 Dr. Brian J. Potter is supported by a *Fonds de recherche du Québec-Santé* career award
365 (267436).

366

367 **Disclosures**

368 Dr. Brian J. Potter has received research funding from Bayer Canada, Boehringer-Ingelheim
369 Canada, and Novartis Canada.

370

371

372

373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425

References

1. Kochanek KD, Xu J, Arias E. Mortality in the United States, 2019. *NCHS Data Brief*. 2020;1-8.
2. Tarride JE, Lim M, DesMeules M, et al. A review of the cost of cardiovascular disease. *Can J Cardiol*. 2009;25:e195-202.
3. Virani SS, Alonso A, Benjamin EJ, et al. Heart Disease and Stroke Statistics-2020 Update: A Report From the American Heart Association. *Circulation*. 2020;141:e139-e596.
4. Arnett DK, Blumenthal RS, Albert MA, et al. 2019 ACC/AHA Guideline on the Primary Prevention of Cardiovascular Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation*. 2019;140:e596-e646.
5. Ivers NM, Jiang M, Alloo J, et al. Diabetes Canada 2018 clinical practice guidelines: Key messages for family physicians caring for patients living with type 2 diabetes. *Can Fam Physician*. 2019;65:14-24.
6. Anderson TJ, Gregoire J, Pearson GJ, et al. 2016 Canadian Cardiovascular Society Guidelines for the Management of Dyslipidemia for the Prevention of Cardiovascular Disease in the Adult. *Can J Cardiol*. 2016;32:1263-1282.
7. Brunham LR, Ruel I, Aljenedil S, et al. Canadian Cardiovascular Society Position Statement on Familial Hypercholesterolemia: Update 2018. *Can J Cardiol*. 2018;34:1553-1563.
8. Nerenberg KA, Zarnke KB, Leung AA, et al. Hypertension Canada's 2018 Guidelines for Diagnosis, Risk Assessment, Prevention, and Treatment of Hypertension in Adults and Children. *Can J Cardiol*. 2018;34:506-525.
9. Ezekowitz JA, O'Meara E, McDonald MA, et al. 2017 Comprehensive Update of the Canadian Cardiovascular Society Guidelines for the Management of Heart Failure. *Can J Cardiol*. 2017;33:1342-1433.
10. Mehta SR, Bailey KR, Cantor WJ, et al. 2018 Canadian Cardiovascular Society/Canadian Association of Interventional Cardiology Focused Update of the Guidelines for the Use of Antiplatelet Therapy. *Can J Cardiol*. 2018;34:214-233.
11. Brass EP, Hiatt WR. Aspirin Monotherapy Should Not Be Recommended for Cardioprotection in Patients With Symptomatic Peripheral Artery Disease. *Circulation*. 2017;136:785-786.
12. Powers WJ, Rabinstein AA, Ackerson T, et al. 2018 Guidelines for the Early Management of Patients With Acute Ischemic Stroke: A Guideline for Healthcare Professionals From the American Heart Association/American Stroke Association. *Stroke*. 2018;49:e46-e110.
13. Cacoub PP, Zeymer U, Limbourg T, et al. Effects of adherence to guidelines for the control of major cardiovascular risk factors on outcomes in the REDuction of Atherothrombosis for Continued Health (REACH) Registry Europe. *Heart*. 2011;97:660-667.
14. Komajda M, Cowie MR, Tavazzi L, et al. Physicians' guideline adherence is associated with better prognosis in outpatients with heart failure with reduced ejection fraction: the QUALIFY international registry. *Eur J Heart Fail*. 2017;19:1414-1423.
15. Nguyen T, Le KK, Cao HTK, et al. Association between in-hospital guideline adherence and postdischarge major adverse outcomes of patients with acute coronary syndrome in Vietnam: a prospective cohort study. *BMJ Open*. 2017;7:e017008.
16. Shah BR, O'Brien EC, Roe MT, Chen AY, Peterson ED. The association of in-hospital guideline adherence and longitudinal postdischarge mortality in older patients with non-ST-segment elevation myocardial infarction. *Am Heart J*. 2015;170:273-280 e271.
17. Avezum A, Oliveira GBF, Lanús F, et al. Secondary CV Prevention in South America in a Community Setting: The PURE Study. *Glob Heart*. 2017;12:305-313.

- 426 18. Eapen ZJ, Liang L, Shubrook JH, et al. Current quality of cardiovascular prevention for
427 Million Hearts: an analysis of 147,038 outpatients from The Guideline Advantage. *Am*
428 *Heart J.* 2014;168:398-404.
- 429 19. Farkouh ME, Boden WE, Bittner V, et al. Risk factor control for coronary artery disease
430 secondary prevention in large randomized trials. *J Am Coll Cardiol.* 2013;61:1607-
431 1615.
- 432 20. Machline-Carrion MJ, Soares RM, Damiani LP, et al. Rationale and design for a cluster
433 randomized quality-improvement trial to increase the uptake of evidence-based
434 therapies for patients at high cardiovascular risk: The BRIDGE-Cardiovascular
435 Prevention trial. *Am Heart J.* 2019;207:40-48.
- 436 21. Solomon MD, Leong TK, Levin E, et al. Cumulative Adherence to Secondary
437 Prevention Guidelines and Mortality After Acute Myocardial Infarction. *J Am Heart*
438 *Assoc.* 2020;9:e014415.
- 439 22. Chow CK, Teo KK, Rangarajan S, et al. Prevalence, awareness, treatment, and control
440 of hypertension in rural and urban communities in high-, middle-, and low-income
441 countries. *JAMA.* 2013;310:959-968.
- 442 23. Schwalm JD, McCready T, Lear SA, et al. Exploring New Models for Cardiovascular
443 Risk Reduction: The Heart Outcomes Prevention and Evaluation 4 (HOPE 4) Canada
444 Pilot Study. *CJC Open.* 2021;3:267-275.
- 445 24. Hosseinzadeh-Shanjani Z, Hoveidamanesh S, Ramezani M, Davoudi F, Nojomi M.
446 Adherence of cardiologist physicians to the American Heart Association guideline in
447 approach to risk factors of cardiovascular diseases: An experience from a teaching
448 hospital. *ARYA Atheroscler.* 2019;15:38-43.
- 449 25. Bhatt DL, Scheiman J, Abraham NS, et al. ACCF/ACG/AHA 2008 expert consensus
450 document on reducing the gastrointestinal risks of antiplatelet therapy and NSAID use:
451 a report of the American College of Cardiology Foundation Task Force on Clinical
452 Expert Consensus Documents. *Circulation.* 2008;118:1894-1909.
- 453 26. Eikelboom JW, Connolly SJ, Bosch J, et al. Rivaroxaban with or without Aspirin in
454 Stable Cardiovascular Disease. *N Engl J Med.* 2017;377:1319-1330.
- 455 27. Hobin E, Shokar S, Vallance K, et al. Communicating risks to drinkers: testing alcohol
456 labels with a cancer warning and national drinking guidelines in Canada. *Can J Public*
457 *Health.* 2020;111:716-725.
- 458 28. von Elm E, Altman DG, Egger M, et al. The Strengthening the Reporting of
459 Observational Studies in Epidemiology (STROBE) Statement: guidelines for reporting
460 observational studies. *Int J Surg.* 2014;12:1495-1499.
- 461 29. Bosch J, Eikelboom JW, Connolly SJ, et al. Rationale, Design and Baseline
462 Characteristics of Participants in the Cardiovascular Outcomes for People Using
463 Anticoagulation Strategies (COMPASS) Trial. *Can J Cardiol.* 2017;33:1027-1035.
- 464 30. Marso SP, Daniels GH, Brown-Frandsen K, et al. Liraglutide and Cardiovascular
465 Outcomes in Type 2 Diabetes. *N Engl J Med.* 2016;375:311-322.
- 466 31. Neal B, Perkovic V, Mahaffey KW, et al. Canagliflozin and Cardiovascular and Renal
467 Events in Type 2 Diabetes. *N Engl J Med.* 2017;377:644-657.
- 468 32. Anker SD, Butler J, Filippatos G, et al. Empagliflozin in Heart Failure with a Preserved
469 Ejection Fraction. *N Engl J Med.* 2021;385:1451-1461.
- 470 33. Anker SD, Butler J, Filippatos GS, et al. Evaluation of the effects of sodium-glucose
471 co-transporter 2 inhibition with empagliflozin on morbidity and mortality in patients with
472 chronic heart failure and a preserved ejection fraction: rationale for and design of the
473 EMPEROR-Preserved Trial. *Eur J Heart Fail.* 2019;21:1279-1287.
- 474 34. McMurray JJV, Docherty KF, Jhund PS. Dapagliflozin in Patients with Heart Failure
475 and Reduced Ejection Fraction. Reply. *N Engl J Med.* 2020;382:973.
- 476 35. Packer M, Anker SD, Butler J, et al. Cardiovascular and Renal Outcomes with
477 Empagliflozin in Heart Failure. *N Engl J Med.* 2020;383:1413-1424.
- 478 36. Perk J, De Backer G, Gohlke H, et al. European Guidelines on cardiovascular disease
479 prevention in clinical practice (version 2012). The Fifth Joint Task Force of the
480 European Society of Cardiology and Other Societies on Cardiovascular Disease

- 481 Prevention in Clinical Practice (constituted by representatives of nine societies and by
482 invited experts). *Eur Heart J.* 2012;33:1635-1701.
- 483 **37.** Mosca L, Linfante AH, Benjamin EJ, et al. National study of physician awareness and
484 adherence to cardiovascular disease prevention guidelines. *Circulation.*
485 2005;111:499-510.
486

487

488

489

490

491

492

493

494

495

496

497

498

499

500

501

502

503

504

505

506

507

508

509

510

511

512

513

514

Tables

515

516 **Table 1.** Characteristics of the total cohort.

Characteristic	Total cohort (N=2503)
Age, mean \pm SD	66 \pm 15
Age >40 years, n (%)	2346 (94)
Male, n (%)	1506 (60)
Weight, mean \pm SD (Kg)	79 \pm 20
BMI \geq 30, n (%)	339 (14)
Missing data, n (%)	1468 (59)
NYHA, median (IQR)	2 (1-2)
Missing data, n (%)	1513 (60)
\geq 2 heart failure hospitalizations in the past year, n (%)	45 (2)
Comorbidities	
Diabetes, n (%)	644 (26)
Dyslipidemia, n (%)	1567 (63)
CAD, n (%)	1032 (41)
PAD, n (%)	318 (13)
HFrEF, n (%)	184 (7)
Hypertension, n (%)	1378 (55)
CKD, n (%)	438 (18)
Lifestyle habits	
Active smoking, n (%)	188 (8)
Alcohol more than recommendations*, n (%)	40 (2)
Physical activity >150 minutes per week, n (%)	83 (3)
BMI: body mass index, CAD: coronary artery disease, Chronic kidney disease (eGFR<60ml/min/m ²), HFrEF: heart failure with reduced ejection fraction, IQR: interquartile range; Kg: kilogram, NYHA: New York Heart Association functional class, PAD: peripheral artery disease, SD: standard deviation. *(Women > 11 drinks per week, Men > 14 drinks per week ²⁷)	

517 **Table 2.** Adherence to prevention guidelines in a cardiology clinic.

Risk factor	Optimal adherence to respective guidelines n (%)
Diabetes Glycemic control	300 (47)
Dyslipidemia N=1567 LDL control	608 (39)
Vascular disease Accurate aspirin therapy CAD PAD	782 (95) 160 (92)
HFrEF Triple therapy	62 (34)
Hypertension BP control	899 (65)
Smoking discussion	172 (91)
Alcohol discussion	36 (90)
Obesity Physical activity discussion Diet discussion	93 (27) 45 (11)
BP: blood pressure, CAD: coronary artery disease, HFrEF: heart failure with reduced ejection fraction, PAD: peripheral artery disease.	

518

519