

# Comparative effectiveness of angiotensin-receptor blockers for preventing macrovascular disease in patients with diabetes: a population-based cohort study

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See related commentary by Cooper on page 1023 and at [www.cmaj.ca/lookup/doi/10.1503/cmaj.130507](http://www.cmaj.ca/lookup/doi/10.1503/cmaj.130507)

## ABSTRACT

**Background:** Telmisartan, unlike other angiotensin-receptor blockers, is a partial agonist of peroxisome proliferator-activated receptor- $\gamma$ , a property that has been associated with improvements in surrogate markers of cardiovascular health in small trials involving patients with diabetes. However, whether this property translates into a reduced risk of cardiovascular events and death in these patients is unknown. We sought to explore the risk of myocardial infarction, stroke and heart failure in patients with diabetes who were taking telmisartan relative to the risk of these events occurring in patients taking other angiotensin-receptor blockers.

**Methods:** We conducted a population-based, retrospective cohort study of Ontario residents with diabetes aged 66 years and older who started treatment with candesartan, irbesartan, losartan, telmisartan or valsartan between Apr. 1, 2001, and Mar. 31, 2011. Our primary outcome was a composite of admission to hospital for acute myocardial infarction, stroke or heart failure. We examined each outcome individually in secondary analyses, in addition to all-cause mortality.

**Results:** We identified 54 186 patients with diabetes who started taking an angiotensin-receptor blocker during the study period. After multivariable adjustment, patients who took either telmisartan (adjusted hazard ratio [HR] 0.85, 95% confidence interval [CI] 0.74–0.97) or valsartan (adjusted HR 0.86, 95% CI 0.77–0.95) had a lower risk of the composite outcome compared with patients who took irbesartan. In contrast, no significant difference in risk was seen between other angiotensin-receptor blockers and irbesartan. In secondary analyses, we found a reduced risk of admission to hospital for heart failure with telmisartan compared with irbesartan (adjusted HR 0.79, 95% CI 0.66–0.96), but no significant differences in risk were seen between angiotensin-receptor blockers in our other secondary analyses.

**Interpretation:** Compared with other angiotensin-receptor blockers, telmisartan and valsartan were both associated with a lower risk of admission to hospital for acute myocardial infarction, stroke or heart failure among older adults with diabetes and hypertension. Telmisartan and valsartan may therefore be the preferred angiotensin-receptor blockers for use in these patients.

**Competing interests:** Tony Antoniou has received unrestricted research grants from Merck and Pfizer. Tara Gomes has received grant funding for the Ontario Drug Policy Research Network from the Ontario Ministry of Health and Long-Term Care. Muhammad Mamdani serves on CMAJ's Editorial Advisory Board (but had no role in the decision to publish this paper), has been on other advisory boards and has received honoraria from AstraZeneca, Bristol-Myers Squibb, Eli Lilly and Company, GlaxoSmithKline, Hoffman-La Roche, Novartis, Novo Nordisk, Merck, Sanofi-Aventis and Pfizer. No other competing interests were declared.

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About 366 million people worldwide live with diabetes, a number that is projected to increase to 552 million by 2030.<sup>1</sup> Because disease-attributable macrovascular complications are the principal causes of death for people with type 2 diabetes, many therapies have the goal of reducing vascular events among these patients.<sup>2,3</sup> Blockade of the renin-angiotensin-aldosterone system with angiotensin-receptor blockers is a commonly used and particularly appealing strategy in this regard, given the multiple mechanisms through which angiotensin II contributes to a heightened risk of diabetes-related macrovascular disease and the superior tolerabil-

ity profile of these drugs relative to angiotensin-converting enzyme (ACE) inhibitors.<sup>4,6</sup>

Although angiotensin-receptor blockers are considered largely interchangeable in clinical practice, evidence from experimental studies and small comparative trials suggest that telmisartan exhibits several pleiotropic properties that distinguish it from other members of this drug class.<sup>7</sup> Most notably, telmisartan is a partial agonist of peroxisome proliferator-activated receptor- $\gamma$  (PPAR $\gamma$ ), a property associated with improvements in surrogate markers of cardiovascular health in small trials involving patients with type 2 diabetes.<sup>8–13</sup> However, whether

telmisartan-mediated activation of PPAR $\gamma$  is associated with a reduced risk of macrovascular events in patients with diabetes relative to angiotensin-receptor blockers that do not share this property is unknown.

We sought to compare the risk of acute myocardial infarction, heart failure and stroke in older patients whose diabetes was also managed with either telmisartan or any of candesartan, irbesartan, losartan and valsartan. We speculated

that, owing to its pleiotropic effects, telmisartan would be associated with a lower risk of macrovascular events in these patients relative to other angiotensin-receptor blockers.

## Methods

### Study design

We conducted a population-based retrospective cohort study involving Ontario residents with

**Table 1 (part 1 of 2):** Baseline characteristics of patients

Characteristic	Angiotensin-receptor blocker, no. (%) <sup>*</sup>				
	Telmisartan (n = 8 182)	Candesartan (n = 10 940)	Irbesartan (n = 12 691)	Losartan (n = 8 411)	Valsartan (n = 13 962)
Age, yr, median (IQR)	73 (69–78)	73 (69–79)	73 (69–78)	73 (69–79)	73 (69–78)
Age group, yr					
66–74	4 929 (60.2)	6 426 (58.7)	7 603 (59.9)	4 909 (58.4)	8 197 (58.7)
75–84	2 778 (34.0)	3 778 (34.5)	4 342 (34.2)	2 946 (35.0)	4 863 (34.8)
≥ 85	475 (5.8)	736 (6.7)	746 (5.9)	556 (6.6)	902 (6.5)
Female sex	4 782 (58.4)	6 579 (60.1)	7 326 (57.7)	5 238 (62.3)	8 466 (60.6)
Duration of diabetes, yr, median (IQR)	6.0 (2.3–10.8)	6.3 (2.4–11)	6.5 (2.4–11.1)	6.2 (2.3–10.5)	6.1 (2.3–10.5)
No admission to hospital	6 272 (76.7)	8 177 (74.7)	9 541 (75.2)	6 351 (75.5)	10 725 (76.8)
Charlson comorbidity index					
0	567 (6.9)	801 (7.3)	860 (6.8)	511 (6.1)	978 (7.0)
1	595 (7.3)	801 (7.3)	940 (7.4)	661 (7.9)	1 000 (7.2)
≥ 2	748 (9.1)	1 161 (10.6)	1 350 (10.6)	888 (10.6)	1 259 (9.0)
Reason for admission to hospital in previous 5 yr					
Congestive heart failure	519 (6.3)	989 (9.0)	957 (7.5)	755 (9.0)	997 (7.1)
Angina	163 (2.0)	234 (2.1)	251 (2.0)	209 (2.5)	261 (1.9)
Diabetes	347 (4.2)	557 (5.1)	686 (5.4)	432 (5.1)	553 (4.0)
Hypertension	2 867 (35.0)	4 234 (38.7)	4 537 (35.8)	3 252 (38.7)	5 167 (37.0)
Stroke or transient ischemic attack	261 (3.2)	417 (3.8)	443 (3.5)	312 (3.7)	431 (3.1)
Hypertrophic cardiomyopathy	11 (0.1)	15 (0.1)	9 (0.1)	9 (0.1)	14 (0.1)
Atrial fibrillation	186 (2.3)	308 (2.8)	303 (2.4)	247 (2.9)	336 (2.4)
Ventricular dysrhythmia	194 (2.4)	332 (3.0)	313 (2.5)	273 (3.3)	353 (2.5)
Peripheral vascular disease	58 (0.7)	100 (0.9)	107 (0.8)	77 (0.9)	96 (0.7)
Percutaneous coronary intervention	907 (11.1)	1 402 (12.8)	1 591 (12.5)	1 091 (13.0)	1 645 (11.8)
Coronary artery bypass grafting	10 (0.1)	6 (0.1)	≤ 5 (0.0)	8 (0.1)	6 (0.0)
History of chronic liver disease in 1 yr	92 (1.1)	153 (1.4)	180 (1.4)	122 (1.5)	210 (1.5)
History of chronic kidney disease in 1 yr	713 (8.7)	1 262 (11.5)	1 430 (11.3)	872 (10.4)	1 148 (8.2)
Residence in a long-term care facility	148 (1.8)	206 (1.9)	243 (1.9)	146 (1.7)	221 (1.6)
No. of prescription drugs in previous 1 yr, median (IQR)	8 (5–12)	9 (6–12)	8 (6–12)	8 (5–12)	8 (5–12)
Medications used in previous 1 yr					
Angiotensin-converting enzyme inhibitors	4 740 (57.9)	6 562 (60.0)	7 833 (61.7)	4 969 (59.1)	8 372 (60.0)
Acetylsalicylic acid	868 (10.6)	1 296 (11.9)	1 436 (11.3)	1 091 (13.0)	1 677 (12.0)
Other antiplatelet drugs	209 (2.6)	326 (3.0)	363 (2.9)	204 (2.4)	332 (2.4)
$\beta$ -adrenergic receptor antagonists	1 953 (23.9)	2 734 (25.0)	3 116 (24.6)	1 996 (23.7)	3 372 (24.2)
Calcium-channel blockers	2 740 (33.5)	3 896 (35.6)	4 649 (36.6)	2 805 (33.4)	4 823 (34.5)

Continued

diabetes aged 66 years and older who started treatment with either candesartan, irbesartan, telmisartan, losartan or valsartan between Apr. 1, 2001, and Mar. 31, 2010.

### Data sources

We determined medication exposure using data from the Ontario Drug Benefit database, which contains comprehensive records of prescription drugs dispensed to Ontario residents aged

65 years and older. We excluded the first year of eligibility for prescription drug coverage (age 65 yr) to avoid having incomplete medication records. We obtained data on hospital admissions from the Canadian Institute for Health Information's Discharge Abstract Database, which contains detailed clinical information regarding all hospital admissions in Ontario. The abstraction of patient charts is undertaken by trained health information professionals using standard diagno-

**Table 1 (part 2 of 2):** Baseline characteristics of patients

Characteristic	Angiotensin-receptor blocker, no. (%)*				
	Telmisartan (n = 8 182)	Candesartan (n = 10 940)	Irbesartan (n = 12 691)	Losartan (n = 8 411)	Valsartan (n = 13 962)
Medications used in previous 1 yr					
Thiazide diuretics	1 842 (22.5)	2 338 (21.4)	2 846 (22.4)	1 810 (21.5)	2 960 (21.2)
Aldosterone antagonists	162 (2.0)	281 (2.6)	277 (2.2)	177 (2.1)	313 (2.2)
Other diuretic agents	1 533 (18.7)	2 256 (20.6)	2 654 (20.9)	1 655 (19.7)	2 727 (19.5)
Other antihypertensive agents	66 (0.8)	107 (1.0)	133 (1.1)	85 (1.0)	140 (1.0)
Nitrates	409 (5.0)	624 (5.7)	708 (5.6)	516 (6.1)	762 (5.5)
Statins	4 411 (53.9)	5 717 (52.3)	6 872 (54.2)	3 957 (47.1)†	7 056 (50.5)
Digoxin	196 (2.4)	320 (2.9)	320 (2.5)	286 (3.4)	405 (2.9)
Amiodarone	46 (0.6)	85 (0.8)	69 (0.5)	58 (0.7)	83 (0.6)
Warfarin	349 (4.3)	593 (5.4)	619 (4.9)	407 (4.8)	629 (4.5)
Nonsteroidal anti-inflammatory drugs	2 257 (27.6)	3 046 (27.8)	3 500 (27.6)	2 282 (27.1)	4 027 (28.8)
Diabetes treatments used in previous 1 yr					
Insulin	709 (8.7)	1 099 (10.1)	1 412 (11.1)	907 (10.8)	1 102 (7.9)
Metformin	3 683 (45.0)	4 906 (44.8)	6 096 (48.0)	3 772 (44.9)	6 526 (46.7)
Acarbose	83 (1.0)	133 (1.2)	151 (1.2)	123 (1.5)	180 (1.3)
Pioglitazone	263 (3.2)	321 (2.9)	368 (2.9)	136 (1.6)	333 (2.4)
Rosiglitazone	213 (2.6)	293 (2.7)	381 (3.0)	197 (2.3)	352 (2.5)
Sulfonylureas	2 237 (27.3)	3 277 (30.0)	4 003 (31.5)	2 758 (32.8)†	4 296 (30.8)
Repaglinide	30 (0.4)	34 (0.3)	59 (0.5)	33 (0.4)	31 (0.2)
Income quintile					
1 (lowest)	1 815 (22.2)	2 555 (23.4)	2 912 (23.0)	1 958 (23.3)	3 213 (23.0)
2	1 798 (22.0)	2 596 (23.7)	2 813 (22.2)	1 908 (22.7)	3 240 (23.2)
3	1 691 (20.7)	2 094 (19.1)	2 534 (20.0)	1 685 (20.0)	2 825 (20.2)
4	1 473 (18.0)	1 941 (17.7)	2 269 (17.9)	1 530 (18.2)	2 536 (18.2)
5 (highest)	1 374 (16.8)	1 722 (15.7)	2 113 (16.6)	1 294 (15.4)	2 112 (15.1)
Missing	31 (0.4)	32 (0.3)	50 (0.4)	36 (0.4)	36 (0.3)
Specialty of prescribing physician					
Family physician/general practitioner	6 376 (77.9)	7 967 (72.8)‡	9 519 (75.0)	5 992 (71.2)†	10 867 (77.8)
Endocrinology	206 (2.5)	247 (2.3)	293 (2.3)	182 (2.2)	172 (1.2)
Nephrology	82 (1.0)	233 (2.1)	302 (2.4)	152 (1.8)	82 (0.6)
Other	632 (7.7)	1 018 (9.3)	996 (7.9)	745 (8.9)	1 032 (7.4)
Unknown	886 (10.8)	1 475 (13.5)	1 581 (12.5)	1 340 (15.9)†	1 809 (13.0)

Note: IQR = interquartile range.

\*Unless otherwise indicated.

†Standardized difference > 0.1 between losartan and telmisartan groups.

‡Standardized difference > 0.1 between candesartan and telmisartan groups.

sis and procedure codes. We used the Ontario Health Insurance Plan database to identify claims for physician services, and the Ontario Diabetes Database to obtain information regarding diabetes diagnoses. This database is a validated administrative data registry of Ontario residents with diagnosed diabetes, which was generated from hospital records and physician services claims. The definition of at least 1 hospital admission or 2 physicians' claims with a diabetes diagnosis within a 2-year period has sensitivity of 86% and specificity of 97% for identifying people with diabetes (excluding gestational diabetes) in primary care records.<sup>14</sup> Finally, we used the Institute for Clinical Evaluative Sciences Physician Database to determine physician specialties and the Registered Persons Database to obtain patient demographic data. These databases were linked in an anonymous fashion using encrypted health card numbers; they are regularly used for population-based drug research.<sup>15-18</sup>

### Identification of cohort

We defined the index date as the date on which the first prescription for a study drug was dispensed. To restrict our analysis to patients taking these drugs for the first time, we excluded people who had received a prescription for any angiotensin-receptor blocker in the year preceding the index date. We also excluded patients who received their diagnosis of diabetes after they were given a pre-

scription for an angiotensin-receptor blocker, patients who had a myocardial infarction in the 5 years preceding the start of treatment and patients who received an ACE inhibitor in conjunction with an angiotensin-receptor blocker.

We considered the use of an angiotensin-receptor blocker to be continuous if a prescription was refilled within 1.5-times the number of days of the preceding prescription's supply.

We included patients who switched between formulations of the same drug, but we censored those who either switched to a different angiotensin-receptor blocker during follow-up or who stopped treatment. We considered treatment to have stopped if a prescription was not refilled within 1.5-times the number of days of the preceding prescription's supply. In addition, we censored patients after 5 years of total observation time, at death or at the end of follow-up (Mar. 31, 2011), whichever occurred first.

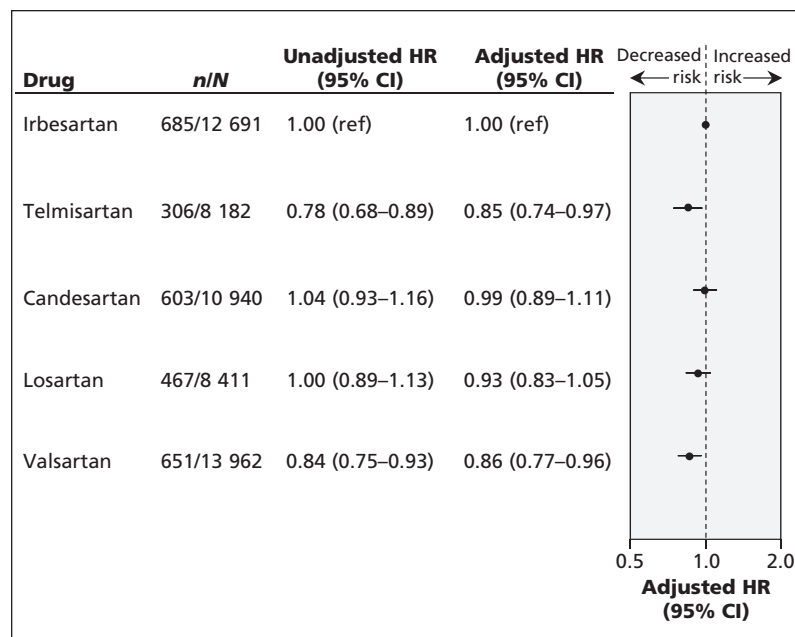
### Outcome measures

Our primary outcome was a composite of admission to hospital for acute myocardial infarction, heart failure or stroke. In secondary analyses, we determined the time to each outcome separately, as well as the time to death from any cause. For patients with multiple admissions to hospital during the study period, we considered only the first admission as an outcome. We determined dates of death using the Registered Persons Database and identified diagnoses during hospital admissions using the relevant codes from the 9th and 10th editions of the International Classification of Diseases (Appendix 1, available at [www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1](http://www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1)).

### Statistical analysis

We calculated descriptive statistics for patients' baseline demographic and clinical characteristics and used standardized differences to test for intergroup differences. Standardized differences of less than 0.1 suggest good balance between groups for a given covariable.<sup>19</sup>

We conducted time-to-event analyses for the primary outcome using multivariable Cox proportional hazards regression to adjust for baseline demographic and clinical variables (Appendix 2, available at [www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1](http://www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1)). We used patients taking irbesartan as the reference group. We selected irbesartan as the referent drug a priori because, like telmisartan, it has a long elimination half-life and has shown superior antihypertensive efficacy relative to comparative angiotensin-receptor blockers.<sup>20-24</sup> Finally, we conducted a dose-response assessment in which we considered low, medium and high doses of angiotensin-receptor



**Figure 1: Risk of the composite outcome (admission to hospital for heart failure, acute myocardial infarction or stroke), by angiotensin-receptor blocker used among older adult patients with diabetes. Ratios were adjusted for all variables listed in Appendix 2 (available at [www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1](http://www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1)). CI = confidence interval, HR = hazard ratio, ref = reference.**

blockers to be time-dependent covariables, using low-dose treatment as the reference. We verified the proportional hazards assumption by testing the statistical significance of a time-dependent treatment variable and by visually inspecting the estimated log(-log) survival curves. All analyses were performed using SAS version 9.2.

This study was approved by the Research Ethics Board of the Sunnybrook Health Sciences Centre, Toronto, Ontario.

## Results

We identified 54 186 patients with diabetes who started treatment with an angiotensin-receptor blocker during the study period. Of these patients, 10 940 (20.2%) took candesartan, 12 691 (23.4%) took irbesartan, 8411 (15.5%) took losartan, 8182 (15.1%) took telmisartan and 13 962 (25.8%) took valsartan (Table 1). For the primary outcome, patients were collectively followed for a total of 107 315 person-years of treatment. Overall, patients were highly similar with respect to demographic characteristics, the specialties of the prescribing physicians, comorbid illnesses and concomitant medications (Table 1).

In the main analysis, the primary outcome (admission to hospital for either acute myocardial infarction, heart failure or stroke) occurred in 2712 (5.0%) patients taking an angiotensin-receptor blocker. After multivariable adjustment (Appendix 2), we found that telmisartan (adjusted hazard ratio [HR] 0.85, 95% confidence interval [CI] 0.74–0.97) and valsartan (adjusted HR 0.86, 95% CI 0.77–0.96) were associated with a significantly lower risk of the primary outcome when compared with irbesartan (Figure 1). In contrast, we found no difference in the risk of the primary outcome between irbesartan and either losartan (adjusted HR 0.93, 95% CI 0.83–1.05) or candesartan (adjusted HR 0.99, 95% CI 0.89–1.11) (Figure 1).

In the secondary analyses, we identified 2708 deaths, 1505 admissions to hospital for heart failure, 806 admissions for acute myocardial infarction and 804 admissions for stroke. We found a lower risk of heart failure with telmisartan versus irbesartan (adjusted HR 0.80, 95% CI 0.66–0.96), but we saw no significant differences between angiotensin-receptor blockers in all other comparisons (Figure 2).

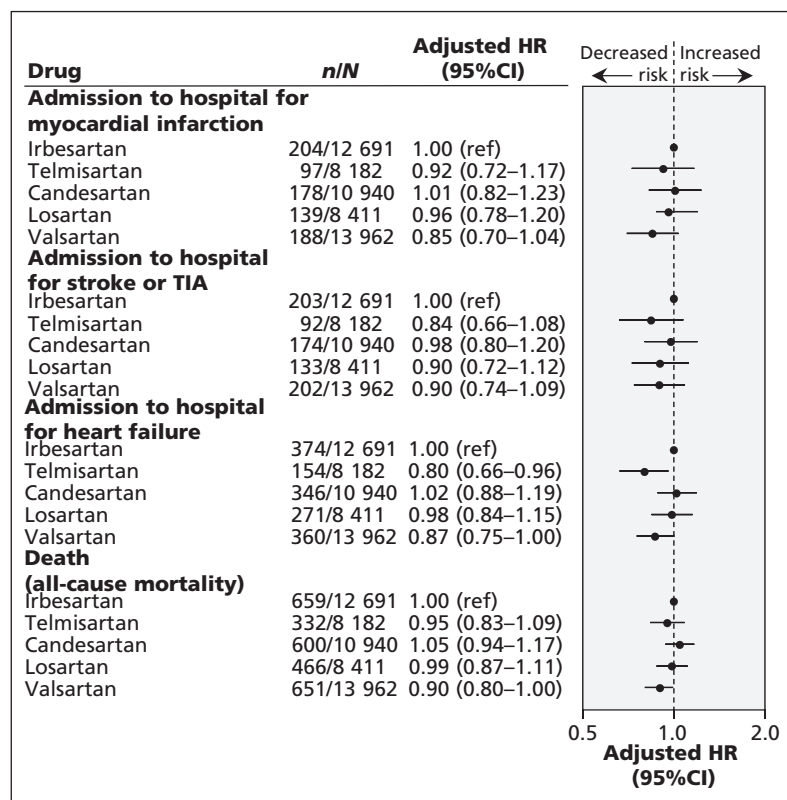
In the dose–response analysis, we found no differences in the risk of the primary outcome with either moderate (adjusted HR 1.04, 95% CI 0.95–1.14) or high (adjusted HR 1.05, 95% CI 0.90–1.23) doses of angiotensin-receptor blockers relative to low doses of these drugs. Our main findings

did not change appreciably after adjusting for dose (Appendix 3, available at [www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1](http://www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1)), although the effectiveness of valsartan was somewhat attenuated (adjusted HR 0.88, 95% CI 0.77–1.0).

## Interpretation

In this population-based study of more than 50 000 older adult patients with diabetes starting treatment with an angiotensin-receptor blocker, we found that valsartan and telmisartan were associated with a reduced risk of admission to hospital for either stroke, myocardial infarction or heart failure relative to irbesartan. Furthermore, telmisartan was associated with a lower risk of admission to hospital for heart failure when compared with irbesartan. Our findings suggest that statistically important differences exist in the effectiveness of angiotensin-receptor blockers when used for the prevention of diabetes-related macrovascular disease, and that a class effect for these agents may not be assumed when used for this purpose in clinical practice.

Although angiotensin-receptor blockers share



**Figure 2:** Risk of individual outcomes (death, or admission to hospital for heart failure, acute myocardial infarction, or stroke), by angiotensin-receptor blockers used among older adult patients with diabetes. Ratios were adjusted for all variables listed in Appendix 2 (available at [www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1](http://www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121771/-/DC1)). CI = confidence interval, HR = hazard ratio, ref = reference, TIA = transient ischemic attack.



common structural features, important pharmacologic differences exist between the drugs that may explain our results. Specifically, at clinically attainable serum concentrations, telmisartan is unique among these drugs in its ability to structurally interact with and activate the PPAR $\gamma$  receptor, a ligand-activated transcription factor that regulates lipid metabolism and insulin sensitivity.<sup>8,25</sup> Because telmisartan is a partial agonist of PPAR $\gamma$ , it is not associated with the adverse effects typically seen with full agonists (e.g., thiazolidinediones), such as sodium and water retention, edema and heart failure.<sup>26,27</sup>

Randomized controlled trials and observational studies comparing cardiovascular outcomes and mortality among patients with diabetes taking individual angiotensin-receptor blockers are lacking. A previous observational study comparing 5 angiotensin-receptor blockers in patients with congestive heart failure found no difference in all-cause mortality between people using telmisartan and those using losartan, but the conclusions were limited by a small sample size (only 143 patients received telmisartan).<sup>28</sup> Our study builds upon the results of smaller trials involving patients with diabetes in which telmisartan was associated with significantly greater reductions in serum lipid levels, plasma glucose concentration, glycated hemoglobin levels and markers of insulin resistance relative to other angiotensin-receptor blockers.<sup>9-13</sup>

In addition, we saw a reduced risk of our primary outcome among patients receiving valsartan. Although this effect was attenuated after adjusting for dose, some evidence suggests that valsartan may impart cardioprotective effects in patients with diabetes unrelated to dose or to the drug's efficacy as an antihypertensive agent. Valsartan has inhibited platelet aggregation in a manner that was neither dose- nor time-dependent, an effect that was more pronounced among patients with diabetes relative to those without the disease.<sup>29</sup>

### Limitations

As with all observational studies, it is possible that our results are biased by intergroup differences in the baseline risk of macrovascular disease and death. However, this seems unlikely, because all groups were highly similar with respect to baseline characteristics, such that any residual differences were negligible and unlikely to account for our results. Furthermore, we adjusted our analyses for an array of important clinical and sociodemographic predictors of angiotensin-receptor blocker use and macrovascular disease.

Although we had no access to clinical information such as smoking history or body mass

index, these limitations apply equally to each of the angiotensin-receptor blockers we investigated. Thus, it is difficult to conceive of an unmeasured variable that would be strongly associated with our outcomes but distributed differentially among the various groups to an extent that could sufficiently compromise our findings. A large randomized controlled trial could conclusively show differences in efficacy among individual angiotensin-receptor blockers in patients with diabetes; however, conducting a trial comparing the efficacy of 5 distinct agents would be expensive and time consuming.

Although miscoding is another possible source of bias in observational studies, we used validated codes for identifying stroke, acute myocardial infection and heart failure in our databases, and differential miscoding among patients using angiotensin-receptor blockers is unlikely.<sup>30-32</sup>

Finally, our study involved patients with diabetes aged 66 years and older. Thus, our findings may not be applicable to younger patients or to patients with hypertension who do not have diabetes.

### Conclusion

Our results suggest that telmisartan and valsartan are associated with a lower risk of admission to hospital for heart failure, stroke or acute myocardial infarction in older patients with diabetes. Pending confirmatory data from additional observational studies or randomized controlled trials, we suggest that a class effect may not be assumed when using angiotensin-receptor blockers for the prevention of diabetes-related macrovascular complications or heart failure, and that telmisartan and valsartan may be the preferred drugs for this indication.

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