

The cost-effectiveness of pharmacist-physician collaborative care models vs usual care on time in target systolic blood pressure range in patients with hypertension: a payer perspective

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What is already known about this subject

- Pharmacist-physician collaborative care models (PPCCMs) are effective at improving blood pressure (BP) control in patients with hypertension, including time in target range (TTR) for systolic BP.
- Increased TTR for systolic BP is associated with reduced risk of adverse cardiovascular events.
- PPCCM for hypertension management is likely cost-effective from the societal perspective when time-based costing methods are used for provider time and BP is studied as a continuous outcome.

What this study adds

- This study assessed the cost-effectiveness of PPCCM from the payer perspective while incorporating the increasingly utilized concept of TTR for systolic BP as a measure of BP control.
- For every 10,000 hypertension patients managed with PPCCM vs usual care over a 3-year time horizon, approximately 27 cardiovascular disease deaths, 29 strokes, 21 nonfatal myocardial infarctions, and 12 incident heart failure diagnoses are expected to be averted.
- Over a 3-year time horizon, PPCCM is expected to not only be less costly to administer but also result in downstream health care savings.

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ABSTRACT

BACKGROUND: Hypertension is highly prevalent in the United States, affecting nearly half of all adults (43%). Studies have shown that pharmacist-physician collaborative care models (PPCCMs) for hypertension management significantly improve blood pressure (BP) control rates and provide consistent control of BP. Time in target range (TTR) for systolic BP is a novel measure of BP control consistency that is independently associated with decreased cardiovascular risk. There is no evidence that observed improvement in TTR for systolic BP with a PPCCM is cost-effective.

OBJECTIVE: To compare the cost-effectiveness of a PPCCM with usual care for the management of hypertension from the payer perspective.

METHODS: We used a decision analytic model with a 3-year time horizon based on published literature and publicly available data. The population consisted of adult patients who had a previous diagnosis of high BP (defined as office-based BP \geq 140/90 mmHg) or were receiving antihypertensive medications. Effectiveness data were drawn from 2 published studies evaluating the effect of PPCCMs (vs usual care) on TTR for systolic BP and the impact of TTR for systolic BP on 4 cardiovascular outcomes (nonfatal

myocardial infarction [MI], stroke, heart failure [HF], and cardiovascular disease [CVD] death). The model incorporated direct medical costs, including both programmatic costs (ie, direct costs for provider time) and downstream health care utilization associated with acute cardiovascular events. One-way sensitivity and threshold analyses examined model robustness.

RESULTS: In base-case analyses, PPCCM hypertension management was associated with lower downstream medical expenditures (difference: $-\$162.86$) and lower total program costs (difference: $-\$108.00$) when compared with usual care. PPCCM was associated with lower downstream medical

expenditures across all parameter ranges tested in the deterministic sensitivity analysis. For every 10,000 hypertension patients managed with PPCCM vs usual care over a 3-year time horizon, approximately 27 CVD deaths, 29 strokes, 21 nonfatal MIs, and 12 incident HF diagnoses are expected to be averted.

CONCLUSIONS: This is the first study to evaluate the cost-effectiveness of PPCCM compared to usual care on TTR for systolic BP in adults with hypertension. PPCCM was less costly to administer and resulted in downstream health care savings and fewer acute cardiovascular events relative to usual care. Although further research is needed to evaluate the long-term costs and outcomes of PPCCM, payer coverage of PPCCM services may prevent future health care costs and improve patient cardiovascular outcomes.

Hypertension is highly prevalent in the United States, affecting nearly half of all adults (43%).¹ Hypertension is defined as having a systolic blood pressure (BP) ≥ 130 or diastolic BP ≥ 80 mmHg and is a major risk factor for ischemic heart disease, heart failure (HF), stroke, chronic kidney disease, and death.^{1,2} Only about a quarter (24%) of adults have their hypertension under control. From 2003-2014, it was estimated that hypertension accounted for \$131 billion per year in US health care costs.³

It has been shown that high BP variability is associated with increased risks of all-cause mortality, coronary heart disease, stroke, and end-stage renal disease.⁴⁻⁶ The concept of time in target range (TTR) for systolic BP is a novel measure of BP variability.⁷ A longitudinal study from 15 Veterans Affairs medical centers categorized TTR for systolic BP into 4 quartiles (0%-25%, 26%-50%, 51%-75%, and 76%-100%) and found an inverse and gradual association between TTR and all-cause mortality.⁷ To determine if TTR for systolic BP had an effect on cardiovascular outcomes, Fatani et al conducted a post hoc analysis of SPRINT (Systolic Blood Pressure Intervention Trial) trial data.^{8,9} In the fully adjusted models, the authors found that for every 1 standard deviation increase in TTR for systolic BP, the risk of a first major adverse cardiovascular event significantly decreased.⁸ This study is consistent with other studies suggesting that greater variability in BP is associated with coronary heart disease, stroke, cardiovascular mortality, and all-cause mortality.⁴⁻⁶

Studies have shown that pharmacists play a key role within primary care settings in managing chronic diseases such as hypertension, and clinical pharmacy services decrease overall health care costs.¹⁰⁻¹² A pharmacist-physician collaborative care model (PPCCM) is a practice model where pharmacists provide medication management for

common primary care conditions, often under a collaborative practice agreement with a physician to adjust medications, and order necessary laboratory tests to monitor drug therapy.¹³ PPCCMs have been shown to not only be successful within an office-based setting,¹³ but even within barbershops and churches.^{14,15} A study by Matzke et al found significant improvements ($P < 0.01$) in hemoglobin, BP, and cholesterol in patients with multiple chronic conditions who were in the PPCCM group compared to those seen by usual care. Additionally, hospitalizations declined within the PPCCM group, which led to an estimated cost savings of \$2,619 per patient.¹⁶ Carter et al have conducted multiple randomized clinical trials to assess the effectiveness of PPCCMs for hypertension management and found that patients treated under a PPCCM achieve significantly better mean BP and overall BP control rates.^{17,18} Recently, a study conducted by Dixon et al investigated the impact of PPCCM on TTR for systolic BP, as defined by the proportion of clinical encounters with systolic BP between 120 and 140 mmHg during a 12-month follow-up period.¹³ The mean TTR for systolic BP was significantly higher among PPCCM patients ($46.2\% \pm 24.3\%$) than patients who received usual care ($24.8\% \pm 27.4\%$) ($P < 0.0001$).¹³ Additionally, a majority of patients in the usual care group had a TTR for systolic BP in the lowest quartile (0%-25%), while PPCCM patients were more likely to have TTR for systolic BP in the highest quartile (76%-100%).¹³

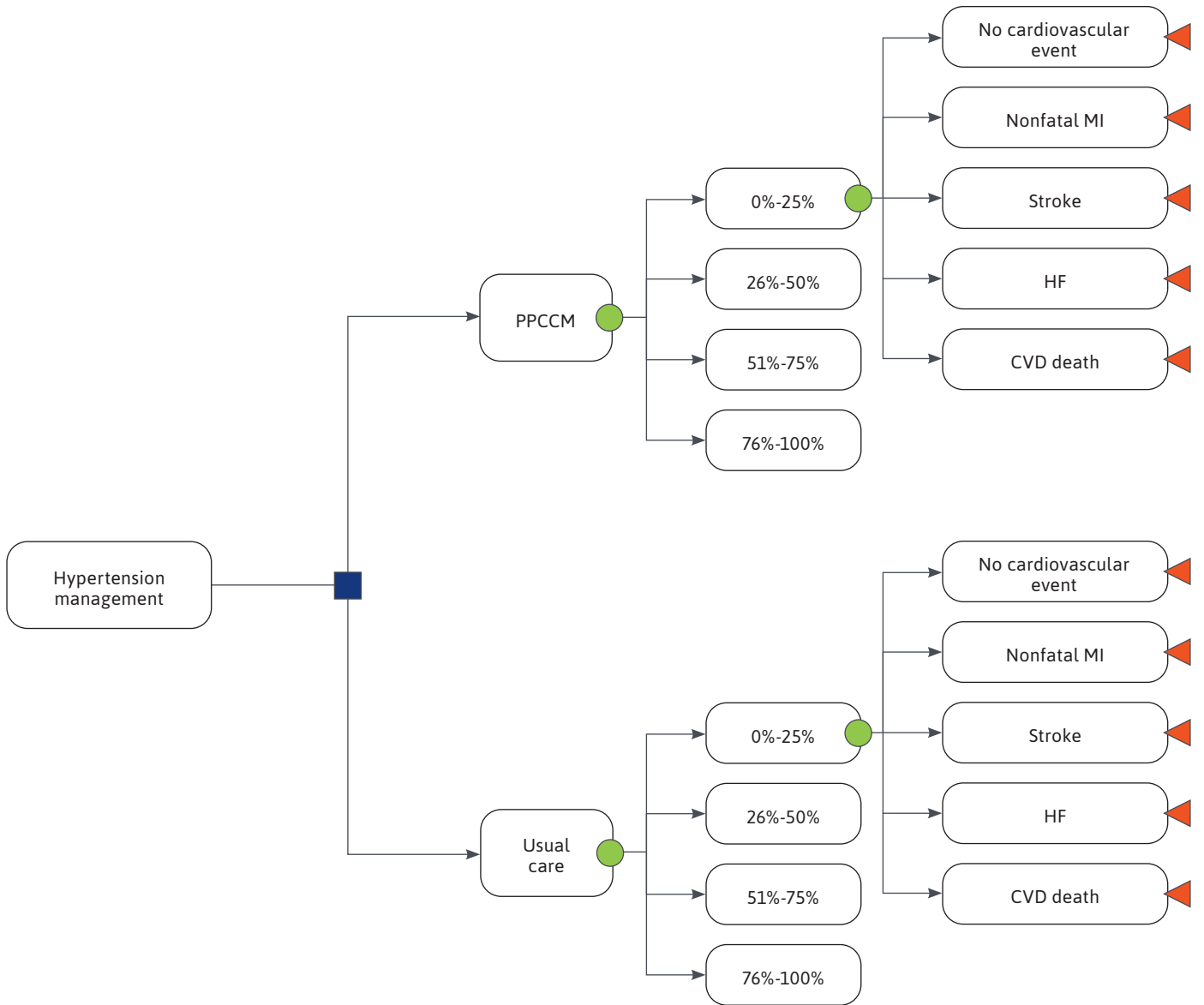
Despite the available evidence supporting PPCCM as an effective model at improving TTR for systolic BP compared to usual care¹³ and that patients with higher TTR for systolic BP have decreased risk of adverse cardiovascular events,⁷⁻⁹ no pharmacoeconomic analysis has combined these findings to model the cost-effectiveness of PPCCM. Therefore, we aimed to compare the cost-effectiveness of PPCCM with usual care on TTR for systolic BP in patients with hypertension. This study was conducted from the payer perspective to quantify the value added to a payer of covering PPCCM services.

Methods

MODEL OVERVIEW

This study used a decision analysis model (Figure 1) to evaluate the cost-effectiveness of 2 hypertension management practices, PPCCM and usual care. The population studied in this analysis consisted of adult patients who were previously diagnosed with hypertension (defined as office-based BP $\geq 140/90$ mmHg) or were receiving antihypertensive medications.¹³ A 3-year time horizon was chosen, reflecting the time frame of available data linking TTR for systolic

FIGURE 1 Decision Tree Analysis for the Cost-Benefit of PPCCM Compared With Standard Usual Care on TTR for Systolic Blood Pressure in Hypertension Management



CVD=cardiovascular disease; HF=heart failure; MI=myocardial infarction; PPCCM=pharmacist-physician collaborative care model; TTR=time in target range.

BP (0%-25%, 26%-50%, 51%-75%, and 76%-100%) to cardiovascular outcome measures (nonfatal myocardial infarction [MI], stroke, HF, and cardiovascular disease [CVD] death).^{8,9} The time horizon is consistent with the follow-up duration from the SPRINT trial, which was terminated early given the clinical benefit of intensive BP control within 3 years

of treatment.^{9,19} Further, the time horizon aligns with the shorter time frame utilized in cost-effectiveness models from the payer perspective.²⁰ The model was developed in TreeAge Pro (TreeAge Software Inc). Institutional review board approval was not required as this research did not qualify as human subject research.

TABLE 1 Effectiveness and Cost Inputs

Variables	Base-case value	Range	Reference
Probability of TTR for systolic BP by hypertension management approach			
PPCCM			
0%-25%	0.210	0.170-0.260	Dixon et al, 2020 ¹³
26%-50%	0.360	0.290-0.430	Dixon et al, 2020 ¹³
51%-75%	0.310	0.240-0.370	Dixon et al, 2020 ¹³
76%-100%	0.120	0.098-0.150	Dixon et al, 2020 ¹³
Usual care			
0%-25%	0.550	0.400-0.600	Dixon et al, 2020 ¹³
26%-50%	0.340	0.270-0.400	Dixon et al, 2020 ¹³
51%-75%	0.050	0.042-0.064	Dixon et al, 2020 ¹³
76%-100%	0.060	0.044-0.066	Dixon et al, 2020 ¹³
Probability of cardiovascular events by TTR for systolic BP			
Outcome event rates of patients in TTR for systolic BP 0%-25%			
Nonfatal MI	0.035	0.027-0.045	Wright et al, 2015 ⁹
Stroke	0.020	0.014-0.028	Wright et al, 2015 ⁹
Heart failure	0.022	0.016-0.031	Wright et al, 2015 ⁹
CVD death	0.017	0.012-0.024	Wright et al, 2015 ⁹
No cardiovascular event	0.906	–	Calculation
Hazard ratio of patients in TTR for systolic BP 26%-50%			
Nonfatal MI	0.83	0.57-1.18	Fatani et al, 2021 ⁸
Stroke	0.83	0.55-1.27	Fatani et al, 2021 ⁸
Heart failure	1.30	0.94-2.01	Fatani et al, 2021 ⁸
CVD death	0.69	0.42-1.15	Fatani et al, 2021 ⁸
No cardiovascular event	1.03	–	Calculation
Hazard ratio of patients in TTR for systolic BP 51%-75%			
Nonfatal MI	0.87	0.61-1.24	Fatani et al, 2021 ⁸
Stroke	0.58	0.36-0.93	Fatani et al, 2021 ⁸
Heart failure	0.84	0.54-1.29	Fatani et al, 2021 ⁸
CVD death	0.53	0.30-0.92	Fatani et al, 2021 ⁸
No cardiovascular event	1.12	–	Calculation
Hazard ratio of patients in TTR for systolic BP 76%-100%			
Nonfatal MI	0.69	0.46-1.04	Fatani et al, 2021 ⁸
Stroke	0.40	0.22-0.73	Fatani et al, 2021 ⁸
Heart failure	0.59	0.34-1.02	Fatani et al, 2021 ⁸
CVD death	0.45	0.23-0.86	Fatani et al, 2021 ⁸
No cardiovascular event	1.25	–	Calculation

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EFFECTIVENESS: TIME IN TARGET SYSTOLIC BP RANGE AND CARDIOVASCULAR EVENT OUTCOMES

Base-case parameters are listed in Table 1. The probabilities that patients managed with PPCCM and usual care would achieve levels of BP control within each of the 4 TTRs for systolic BP quartiles were based on previously published data.¹³ Although published data on the effectiveness of PPCCM had a 1-year study duration,¹³ subjects in the model were assumed to stay in the same quartile of target BP range over the 3-year time horizon to facilitate linking the PPCCM effectiveness data to the clinical data on the association between TTR for systolic BP and cardiovascular events.

Four cardiovascular events (non-fatal MI, stroke, HF, and CVD death) were selected for model inclusion based on available probabilities and hazard ratios from published data on cardiovascular outcomes associated with TTR for systolic BP quartiles.^{8,9} Specifically, data on TTR for systolic BP quartiles and cardiovascular outcomes were derived from a post hoc analysis of the SPRINT trial, a randomized, controlled, open-label trial of intensive vs standard BP control.⁹

Patients within the SPRINT trial were censored after their first cardiovascular event, precluding analysis of subsequent events. Accordingly, cardiovascular events in this study were assumed to be mutually exclusive. Patients who did not incur 1 of these 4 events were assumed to have had no major cardiovascular event.

HYPERTENSION MANAGEMENT AND CARDIOVASCULAR EVENT COSTS

The model incorporated direct medical costs, including both programmatic costs (ie, direct costs for provider time) and downstream health care utilization associated

TABLE 1 Effectiveness and Cost Inputs (continued)

Variables	Base-case value	Range	Reference
Programmatic costs			
Annual PPCCM pharmacist visits, n	6	4-12	Dixon et al, 2020 ¹³
PPCCM cost per visit, \$	24	19-29	ASHP, 2019 ²¹
Annual physician visits			
PPCCM group, n	1	1-2	Assumption
Usual care visits, n	3	1-6	Dixon et al, 2020 ¹³
Physician cost per visit, \$	90	72-108	CMS, 2019 ²²
Total cost of PPCCM, \$	702	562-842	ASHP, 2019 ²¹
Total cost of usual care, \$	810	648-972	CMS, 2019 ²²
Downstream health care costs, \$			
One-time cost of nonfatal MI	24,089	15,372-32,306	Bress et al, 2017 ¹⁹
One-time cost of stroke	15,678	6,001-42,039	Bress et al, 2017 ¹⁹
One-time cost of heart failure	11,678	11,669-16,580	Bress et al, 2017 ¹⁹
One-time cost of CVD death	19,514	12,560-33,024	Bress et al, 2017 ¹⁹

ASHP=American Society for Health-Systems Pharmacists; BP=blood pressure; CMS=Centers for Medicare & Medicaid Services; CVD=cardiovascular disease; MI=myocardial infarction; PPCCM=pharmacist-physician collaborative care model; TTR=time in target range.

with cardiovascular events (Table 1). Cost data were obtained from publicly available data and recently published cost-effectiveness analyses.^{19,21,22}

Provider visit utilization data were obtained from a real-world analysis of PPCCM vs usual care for the management of hypertension.¹³ Specifically, for the cost of the PPCCM program, patients were assumed to have been seen for hypertension management 6 times per year by a pharmacist¹³ and once per year by a physician. Subjects in the usual care group were assumed to be seen 3 times per year by a physician.¹³

The cost per pharmacist visit reflected Current Procedural Terminology (CPT) code 99211 (level 1), an “incident-to” billing code used by pharmacists given a lack of provider status and eligibility to bill at a higher level.²¹ For usual care visits, the CPT code 99213 was used for evaluation and management/outpatient visits.²²

One-time costs of treating each cardiovascular event were obtained from the cost-effectiveness analysis of the SPRINT trial.¹⁹

Costs of hypertensive medications were assumed to be the same for both PPCCM and usual care given a lack of comparative medication use data and hence excluded from the model. Additionally, since the most commonly utilized hypertensive medications are generic and typically inexpensive,²³ they were unlikely to have a major impact on costs of care.

All costs were inflated to 2020 US dollars using the medical care component of the Consumer Price Index.

SENSITIVITY ANALYSES

One-way deterministic sensitivity analyses were performed on all model variables to account for uncertainty in the parameter estimates for the 2 hypertensive management options. Additionally, threshold analyses were

performed varying the cost per pharmacist visit, the number of annual pharmacist visits among patients in the PPCCM program, and the number of annual physician visits among patients in usual care to assess the values at which the programmatic costs of the 2 models would be equal.

Results

In base-case analyses, PPCCM hypertension management was associated with lower total program costs (difference: −\$108.00) and lower downstream medical expenditures (difference: −\$162.86) when compared to usual care (Table 2). For every 10,000 hypertension patients managed with PPCCM vs usual care over a 3-year time horizon, approximately 27 CVD deaths, 29 strokes, 21 nonfatal MIs, and 12 incident HF diagnoses are expected to be averted.

PPCCM was associated with lower downstream medical expenditures across all parameter ranges tested in the deterministic sensitivity analysis. The expected downstream health care savings were most sensitive to the likelihood that patients receiving usual care spend little to no time in therapeutic systolic BP range (TTR for systolic BP: 0%-25%; Figure 2). PPCCM was expected to reduce health care expenditures even as the proportion of usual care patients with TTR for systolic BP of 0%-25% was varied from its base-case value of 55%, the probability observed by Dixon et al,¹³ to the lowest probability tested, 40%.

The program costs of hypertension management with PPCCM, while lower than those of usual care in base-case analyses, were sensitive to the number of visits with a physician (usual care patients) and pharmacist (PPCCM patients; Figure 3). Due to the substantial difference in CPT code reimbursement for pharmacist vs usual care visits, a patient in

TABLE 2 Cost-Effectiveness Results

	PPCCM	Usual care	Difference
Cardiovascular events			
Nonfatal MI	0.0300	0.0321	21 per 10,000
Stroke	0.0149	0.0178	29 per 10,000
Heart failure	0.0225	0.0237	12 per 10,000
CVD death	0.0116	0.0143	27 per 10,000
Total downstream health care expenditures, \$	1,535.82	1,698.64	-\$162.82
Total program costs, \$	702.00	810.00	-\$108.00
Cost-benefit ratio	Dominant		

CVD=cardiovascular disease; MI=myocardial infarction; PPCCM=pharmacist-physician collaborative care model.

the PPCCM program who was seen 6 times per year by a pharmacist and once per year by a physician was still cheaper than a patient in the usual care group who was seen 3 times per year by a physician.

However, in 1-way sensitivity analysis, the cost of PPCCM hypertension management exceeded the cost of usual care when independently varying the number of both types of provider visits. First, if the number of hypertension-related physician visits each year was reduced from 3 to 1 while holding the number of PPCCM-related visits constant, the cost of the PPCCM hypertension management exceeded the cost of usual care by \$432 over the 3-year study period. Second, when the number of pharmacist visits among patients enrolled in PPCCM increased from 6 per year to 12 while the number of physician visits in the usual care group (n=3) was held constant, PPCCM was associated with an incremental program cost of \$324 over usual care.

In a threshold analysis, the costs of the PPCCM and usual care programs became equal when the unit cost of pharmacist visits were increased 62.5%, to \$39. The program costs were also equal when the number of PPCCM patient visits increased from

6 to 10 pharmacist visits per year, or the number of usual care patient visits decreased from 3 to 2 physician visits per year.

Discussion

This study quantifies the cost-effectiveness of PPCCM for hypertension management to improve BP control and cardiovascular outcomes. Previous studies evaluated PPCCM impact on TTR for systolic BP¹³ and the association between TTR for systolic BP and cardiovascular outcomes,⁸ but no pharmacoeconomic analysis had combined these findings to model the cost-effectiveness of PPCCM from the payer perspective.

This study found that patients enrolled in the PPCCM incurred fewer costs associated with their direct hypertension management. The lower PPCCM program costs reflect the significantly lower cost of pharmacist time as billed by “incident to” CPT codes than physician visits for hypertension.

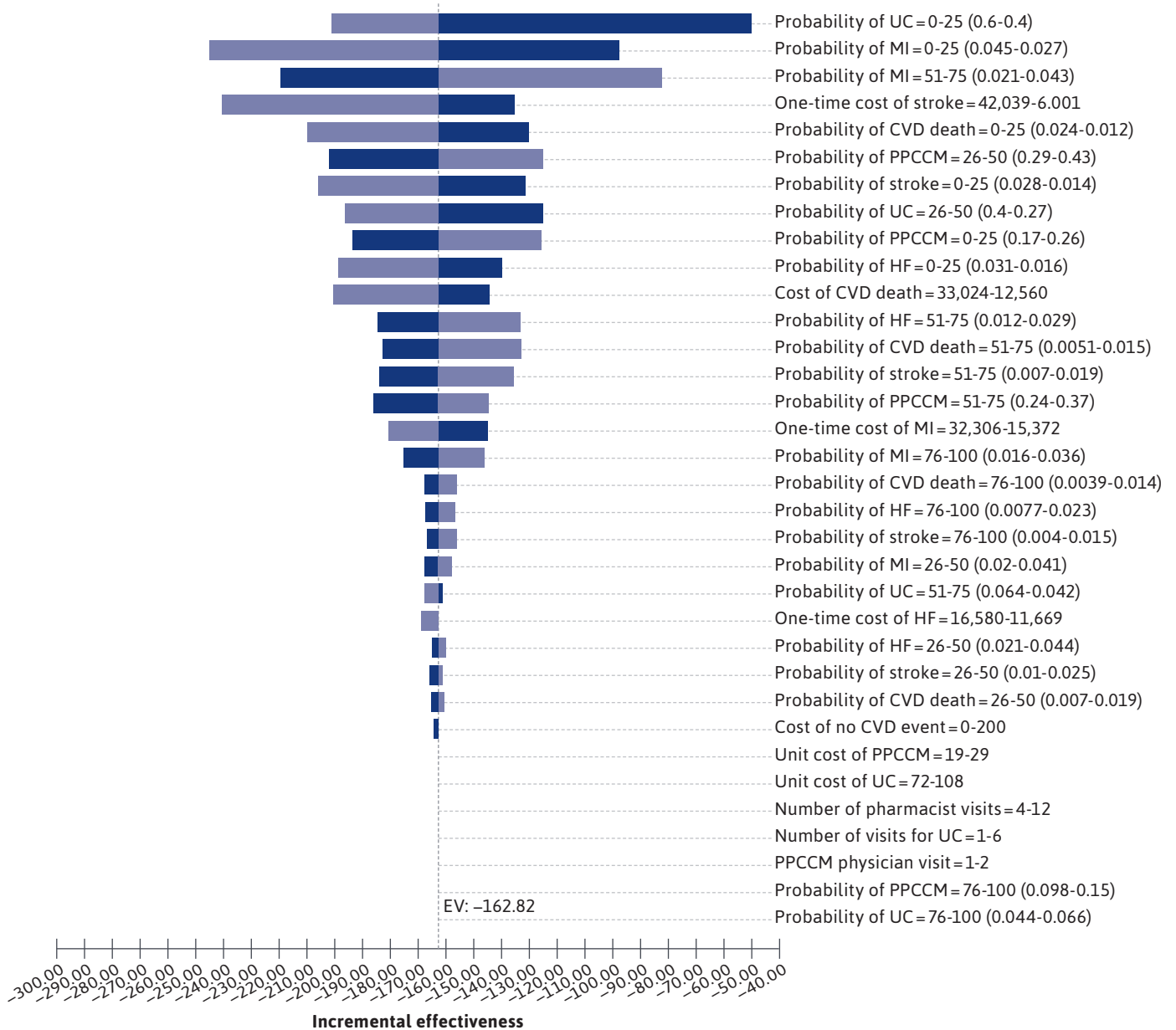
In a threshold analysis, the direct cost of provider time was lower for usual care if patients receiving usual care had fewer than 2 physician visits per year. However, previous studies

suggest that approximately 80% of adult patients with hypertension have 2 or more hypertension-focused physician visits per year.²⁴ Nonetheless, given that the cost of PPCCM hypertension management exceeded the cost of usual care among patients with only 1 hypertension-related physician visit each year, payers concerned with the immediate budget impact of PPCCM reimbursement may focus on coverage for patients with at least 2 or 3 hypertension-related physician visits annually, as PPCCM is cost-neutral and cost-saving, respectively, in these populations.

A second threshold analysis found that the direct program cost of PPCCM would equal that of usual care if patients met 10 times with a pharmacist annually. This well exceeds the number of previously observed pharmacist appointments for patients in 2 different PPCCM programs,^{13,25} suggesting that the PPCCM model is likely to save upfront hypertension management costs from the payer perspective.

While this study found that the direct intervention costs of the PPCCM were lower than those of usual care, several previous cost-effectiveness analyses on pharmacist-physician collaborative care for the management of hypertension found increased costs for patients in a PPCCM.^{25,26} A cost-effectiveness analysis from a societal perspective on a physician-pharmacist collaboration to improve hypertension control conducted by Polgreen et al reported that provider costs over a 9-month period were \$238.96 for PPCCM patients and \$113.67 for usual care patients managed only by a physician.²⁵ Rather than using CPT billing codes, that study determined costs based on time spent with pharmacists and providers and their average compensation rates, likely due to its societal, rather than

FIGURE 2 Tornado Diagram of Incremental Downstream Health Care Expenditures Among Patients Receiving PPCCM vs Usual Care

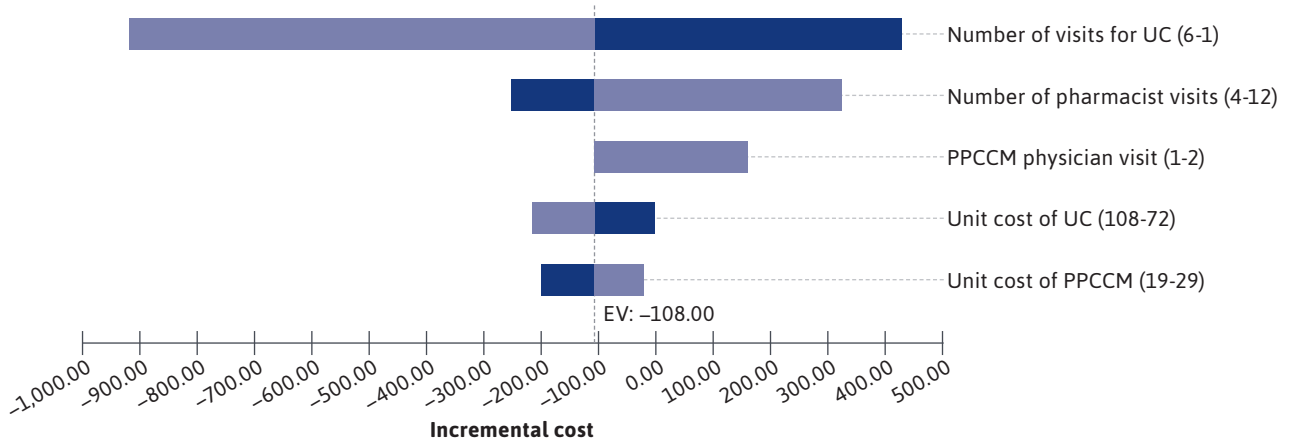


CVD = cardiovascular disease; EV = expected value; HF = heart failure; MI = myocardial infarction; PPCCM = pharmacist-physician collaborative care model; UC = usual care.

payer, perspective. Thus, while usual care patients had the same number of physician visits (median: 3 visits), as was assumed for this analysis, the cost of those 3 visits was calculated to be only \$113.67. Kulchaitanaroaj et al similarly

reported higher costs for PPCCM in 2 analyses,^{26,27} but, like Polgreen,²⁵ used time-based costing, resulting in higher provider costs among PPCCM patients (\$345.25) than those in usual care (\$111.84).²⁷

FIGURE 3 Tornado Diagram of Incremental Cost of PPCCM vs Usual Care



EV = expected value; PPCCM = pharmacist-physician collaborative care model; UC = usual care.

The use of CPT codes in this analysis generated higher expected costs for physician visits but more accurately reflects hypertension management costs from the payer perspective. CPT “incident-to” billing, in which the physician bills, receives payment, and reimburses the pharmacist, offers payers an opportunity to implement payment for services within existing frameworks of physician reimbursement. The use of incident-to billing and collaborative practice agreements may also reduce barriers to PPCCM implementation from the pharmacy perspective, as a lack of clear reimbursement was cited by study authors as a potential barrier to more widespread PPCCM dissemination. Physician champions for the model can help to facilitate reimbursement efforts and streamline referrals, as the pharmacists practicing under the collaborative practice agreement in the collaborative care model routinely reported encountering new complaints from patients.

When compared with usual care, PPCCM was associated with lower downstream health care expenditures, saving an expected \$162.82 over a 3-year time horizon. Our finding of downstream health care savings is consistent with the majority of economic evaluations of clinical pharmacy services for chronic disease state management that incorporate long-term health care expenditures.²⁸ Pharmacist-delivered medication management and hypertension education have consistently been shown to reduce BP,²⁹ which, in turn, is associated with fewer cardiovascular events. Further, the more frequent pharmacist interactions in the PPCCM model may have facilitated the development of a stronger patient-pharmacist relationship and higher

levels of trust, thereby enabling patients to better manage their chronic diseases.³⁰

Our study reported that, for every 10,000 patients with hypertension managed in a PPCCM model, 27 CVD deaths, 29 strokes, 21 nonfatal MIs, and 12 incident HF diagnoses are expected to be averted. While quality-adjusted life-years (QALYs) were not utilized as an outcome in this study, past cost-utility analyses of pharmacist-led or collaborative hypertension management have reported such programs to be cost-effective. Bryant et al modeled 10-year health outcomes and 1-year health care costs associated with pharmacist-led hypertension care in Black-owned barbershops in the Los Angeles Barbershop Blood Pressure Study (LABBPS) from a health care sector perspective.¹⁴ They reported a mean cost of \$42,717 per QALY gained. Kulchaitanaroaj et al similarly reported a PPCCM to be highly cost-effective from the payer perspective (\$26,807 per QALY gained).

This study thus adds to a growing body of literature suggesting that pharmacist collaboration in the management of chronic conditions not only benefits the health outcomes of the patient but does so in a cost-effective manner.³¹⁻³³ PPCCMs may have other benefits not captured in economic evaluations, including decreased physician workload and an ability to reach underserved populations.³⁴

FUTURE RESEARCH

This research reports the cost-benefit of PPCCMs vs usual care on TTR for systolic BP for 4 cardiovascular outcomes. The data for TTR for systolic BP and cardiovascular outcomes

was from previously published data. The direct effect of PPCCM and usual care as it relates to patient outcomes and costs has not been reported. Our study was designed to evaluate only the first occurrence of CVD or death. Therefore, future research that includes a Markov model investigating recurrent cardiovascular events should be conducted. Additionally, the cost-benefit of PPCCM with the addition of hypertensive medication costs should also be explored. Different cardiovascular outcomes can result in additional hypertensive medications, which can affect costs from the payer perspective.

LIMITATIONS

This research included several limitations. TTR for systolic BP data was collected from a study with a small population of 112 patients (56 patients in both PPCCM and usual care), which may limit generalizability,¹³ though the impact of the PPCCM on hypertension management reported by Dixon et al¹³ was similar to that reported elsewhere.^{17,18}

This study did not incorporate the cost of medications due to the lack of information on medication utilization among patients in the 2 groups. The post hoc analysis of SPRINT data by Fatani et al indicated the number of BP-lowering agents based on participants' TTR for systolic BP,⁸ but it is not known whether pharmacist involvement to promote higher TTR for systolic BP would systematically change the number of BP-lowering agents required to improve BP control.

While Dixon et al did not report specific medication utilization in the PPCCM and usual care groups,¹³ the antihypertensives used by both PPCCM and usual care patients were predominately low-cost generics, minimizing the effect of drug costs on the cost-effectiveness of the

program. Additionally, indirect costs were not included in our analysis due to a lack of data linking TTR for systolic BP to changes in productivity, absenteeism, and other indirect costs; similarly, utility values have not yet been established by TTR for systolic BP ranges. However, given that numerous adverse cardiovascular outcomes have been associated with indirect costs of lost productivity due to morbidity and mortality,³⁵ it is likely that the lack of indirect costs in this study resulted in an underestimation of the downstream savings associated with PPCCMs.

Future research is needed to assess indirect costs and potential changes in QALYs associated with improvements in TTR for systolic BP. Furthermore, costs for payer oversight, including quality assurance/auditing of the benefit, were not considered; substantial quality assurance costs may reduce the reported savings associated with PPCCM implementation.

This study evaluated the impact of hypertension management with PPCCM on cardiovascular outcomes and associated costs over a 3-year time frame. Hypertension is a chronic disease and is linked to health consequences including multiple MIs, strokes, and HF exacerbations. The data we utilized from the SPRINT trial censored patients after their first occurrence of a major adverse cardiovascular event during the study period. Therefore, our study may have underestimated the impact of PPCCM on long-term adverse cardiovascular events associated with TTR for systolic BP in hypertension management.

Finally, this study used effectiveness estimates from a real-world study on PPCCM for hypertension management.¹³ However, a recent nationwide survey found that only about half of patients considered themselves likely to participate in clinical pharmacy services under a collaborative practice

agreement, despite their perceptions that such services improve physician-pharmacist coordination.³⁶ If eligible patients choose not to participate in PPCCM services where available, the scope of downstream benefits realized by widespread programmatic access would be more limited than with widespread adoption.

Conclusions

This is the first study to evaluate the cost-benefit of a PPCCM and usual care on TTR for systolic BP in patients with hypertension. The results indicated that a PPCCM was less costly to administer and resulted in reduced downstream adverse cardiovascular events and health care savings relative to usual care. Although further research is needed to evaluate the long-term costs and outcomes of PPCCM, payer coverage of PPCCM services may prevent future health care costs and improve patient cardiovascular outcomes.

DISCLOSURES

No funding was received for the completion of this research. The authors have nothing to disclose.

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