Behavioral Interventions to Improve Hypertension Control in the Veterans Affairs Healthcare System

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Hypertension is a common and costly disease among US veterans. The Veterans Affairs (VA) healthcare system is the largest integrated healthcare provider in the United States and reviewing hypertension interventions developed in the VA may inform interventions delivered in other integrated healthcare systems. This review describes behavioral interventions to improve hypertension control that have been conducted in the VA since 1970. The authors identified 27 articles representing 15 behavioral interventional trials.

In recent decades there has been improvement in hypertension care provided in the Veterans Affairs (VA) healthcare system, the largest integrated healthcare provider in the United States. From 2000 to 2010, blood pressure (BP) control improved in the VA from 43.0% to 76.6%.¹ There is a need for innovative behavioral approaches to continue and improve BP control. Reviewing hypertension interventions developed in the VA may inform interventions delivered in other systems. We performed a literature review describing behavioral hypertension interventions administered in the VA and distilled commonalities among successful programs. We emphasize behavioral management interventions because the success of BP control ultimately depends on a patient's willingness and ability to modify and maintain certain behaviors (eg, proper diet, exercise, and medication adherence). Therefore, our objective was not to report on VA hypertension care in general, but, rather, to focus on behavioral hypertension management interventions.

METHODS

A literature search using the PubMed database was conducted. We identified articles containing MeSH and key words addressing intervention studies or clinical trials, veterans, and hypertension. Articles were limited to those published in English in the past 44 years. Studies need not be solely conducted in the VA. This initial search yielded 171 articles. We screened full

Manuscript received: May 7, 2014; revised: August 21, 2014; accepted: August 24, 2014 DOI: 10.1111/jch.12423 Studies were heterogeneous across patients, providers, interventionist, and intervention components. The VA bridges services related to diagnosis, treatment, medication management, and behavioral counseling in a unified approach that supports collaboration and provides infrastructure for hypertension management. *J Clin Hypertens (Greenwich).* 2014;16:827–837. Published 2014. This article is a U.S. Government work and is in the public domain in the USA.

articles for eligibility. Exclusion criteria included: (1) commentary or editorial; (2) observational or retrospective analysis; (3) review article or solely described a conceptual mode; (4) not focused on hypertension; or (5) not a patient-focused behavioral hypertension intervention. The Figure outlines the article identification process. Interventions were divided into only behavioral or behavioral/medication management. We made this distinction because medication management interventions generally require complex design, necessitating the involvement and/or oversight of a clinician or pharmacist.

RESULTS

We identified 27 articles representing 15 unique trials. Approximately 20% (n=3) of the interventions addressed patients and providers²⁻¹¹ and 80% (n=12) focused on patients solely.¹²⁻²⁷ All studies included patient-directed education; 60% (n=9) of the studies^{6,7,12,14,18-20,23-25} also involved medication management.

Medication Management+Behavioral Components

A 6-month, single-site study was conducted to determine whether pharmaceutical care provided in a pharmacist-managed hypertension clinic or in a traditional primary care setting resulted in better control of hypertension (Table I).²⁶ Individuals in the intervention group were scheduled for a clinical pharmacist visit once monthly. The pharmacist provided drug counseling, addressed recommended lifestyle changes and medication adherence, and made changes in drug selection and dosage as needed. Mean changes in systolic BP from baseline for the intervention and control groups were -18.4 (95% confidence interval [CI], -26.3 to -10.5) and -3.98 (95% CI, -11.8 to

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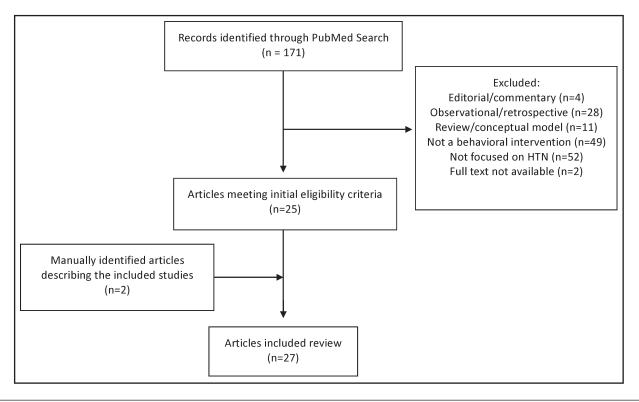


FIGURE. Identification of included articles. The specific search strategy used was: ((((("Hypertension "[Mesh] OR "hypertension" [Title/ Abstract]) AND ("United States Department of Veterans Affairs" [Mesh] OR "Veterans" [Mesh] OR "Veterans Health" [Mesh] OR "Veterans" [Title/Abstract])) AND ("Intervention Studies" [Mesh] OR "Clinical Trial" [Publication Type]) AND English [lang])) AND ("1970/01/01"[PDat]: "2014/ 04/10"[PDat]))).

3.79), respectively (P=.01). There was no significant difference in adherence between (P>.25) or within (P>.07) the two groups at baseline or at the end of the study.²⁶

The Improving Blood Pressure in Colorado (Colorado) study was a 6-month multimodal intervention comprised of patient education, home-based BP monitoring and reporting to an interactive voice-response telephone system, and clinical pharmacist follow-up.¹⁴ The intervention group was given an educational booklet and were trained in home monitoring and instructed to measure their BP three to four times weekly. Pharmacists provided counseling on lifestyle changes and made medication adjustments for those with poor BP control. BP reductions were greater in the intervention vs the usual care group (-13.1 mm Hg vs -7.1 mm Hg, *P*=.006 for systolic; -6.5 mm Hg vs -4.2 mm Hg, *P*=.07 for diastolic).¹⁴

Another intervention involving self-monitoring and medication management was the Hypertension Intervention Nurse Telemedicine Study (HINTS). HINTS evaluated three telephone-based interventions in a fourgroup design: (1) nurse-administered, behavioral management; (2) nurse-administered, physician-directed medication intervention using a validated clinical decision support system; (3) combined behavioral management and medication management intervention; and (4) usual care.¹² All intervention patients were provided with a wireless home BP monitor and advised to monitor their BP daily. Both the behavioral management and medication management alone showed significant improvements at 12 months: 12.8% (95% CI, 1.6%-24.1%) and 12.5% (95% CI, 1.3%-23.6%), respectively. Improvements were not sustained at 18 months. In subgroup analyses, among those with poor baseline BP control, systolic BP decreased in the combined intervention group by 14.8 mm Hg (95% CI, -21.8 mm Hg to -7.8 mm Hg) at 12 months and 8.0 mm Hg (95% CI, -15.5 mm Hg to -0.5 mm Hg) at 18 months, relative to usual care.¹²

A single-site, randomized, controlled device effectiveness study enrolled patients with stage 3 or higher chronic kidney disease and uncontrolled hypertension.²⁴ Patients were randomized to usual care or a telemonitoring device pairing a Bluetooth-enabled BP cuff with an Internetenabled hub, which wirelessly transmitted BP values. For the intervention group, patients were provided with a BP monitor and instructed to follow their physician's instructions regarding the frequency of monitoring. Patients were contacted if BP was above goal. Both groups had a significant improvement in systolic BP (P<.05). Systolic BP fell a median of 13 mm Hg in monitored participants (P for comparison .31).²⁴

TABLE I. Sur Components	mmary of Publi	shed Studies	Describing VA Hype	ertension Control Int	erventions That I	Summary of Published Studies Describing VA Hypertension Control Interventions That Use Medication Management±Behavioral nts	agement±Behaviora	al
Study Title	Lead Investigator (Primary Year)	Study Design	Sample Size	Setting	Ams	Intervention Content	Primary Study Outcome	Duration
Hypertension Intervention Nurse Telemedicine Study (HINTS) ^{12,13,17}	Bosworth (2011)	4-arm randomized controlled trial	591 patients (147 usual care; 148 behavioral management; 149 medication management, 147 combined intervention)	Internal medicine clinics associated with the Durham VA Medical Center Medical Center	Patients were randomized to a usual care control group or 1 of 3 telephone- based intervention groups: nurse- administered behavioral management; nurse- and physician- administered medication management, or a combination of both	Wireless home BP monitoring; behavioral management intervention consisting of 11 tailored health behavioral models; medication management	Change in BP control measured at 6- month intervals	18 months
Veterans Affairs Multi- disciplinary Education and Diabetes Intervention for Cardiac- Extended (MEDIC-E) ²⁰	Cohen (2011)	Randomized controlled trial	99 patients (50 intervention; 49 usual care)	Providence VA Medical Center	Diabetic patients were randomized to a pharmacist- led shared medication appointments program or standard primary care	4 weekly group sessions followed by 5 monthly booster group sessions; each 2-hour session included 1 hour of multidisciplinary diabetes-specific healthy lifestyle education and 1 hour of pharmacotherapeutic intervention performed by a pharmacist	Hemoglobin A _{1 c} , LDL cholesterol, BP, diabetes self-care behavior questionnaire	6 months
N/A ²³	Edelman (2010)	Randomized controlled trial	239 patients were randomized (133 intervention group; 106 usual care group)	Three VA medical centers in North Carolina and Virginia	Patients were randomly assigned within each center to either attend GMCs or receive usual care	GMCs were comprised of 7 or 8 patients and a multidisciplinary care tearn. Groups met every 2 months (7 visits total). At each visit, BP was checked and home blood glucose values were	Outcomes were hemoglobin A _{1c} level and systolic BP	12 months

TABLE I. Summary of Pu Components (Continued)	mmary of Publi: (Continued)	shed Studies I	Describing VA Hyp	bertension Control In	terventions That I	Summary of Published Studies Describing VA Hypertension Control Interventions That Use Medication Management≟Behavioral nts (Continued)	agement±Behavio	ral
Study Title	Lead Investigator (Primary Year)	Study Design	Sample Size	Setting	Arms	Intervention Content	Primary Study Outcome	Duration
The Adherence and Intensification of Medications (AIM) study ^{6,7}	Heisler (2010)	Cluster- randomized controlled effectiveness study	4100 patients (1797 intervention; 2303 control)	Three VA facilities and 2 Kaiser Permanente Northern California facilities	Primary care teams within sites were randomized to a program led by a clinical pharmacist trained in motivational interviewing- based behavioral counseling approaches and authorized to make BP medication cusual care	collated. Patients attended an educational session delivered by the nurse or educator. Topics of the educational sessions were tailored to members' needs. The pharmacist and the primary care internist reviewed patient medical records, BPs, and home blood glucose readings during each session and developed individualized plans for medication or lifestyle management directed toward improving BP and hemoglobin A ₁ c level. Sessions lasted 90 to 120 minutes Motivational interviewing, medication management, behavioral change	Relative change in systolic BP measurements of time	14 months

TABLE I. Summary of P(Components (Continued)	mmary of Publi (Continued)	shed Studies	Describing VA Hype	rtension Control Int	terventions That	Summary of Published Studies Describing VA Hypertension Control Interventions That Use Medication Management±Behavioral nts (Continued)	agement±Behavior	al
Study Title	Lead Investigator (Primary Year)	Study Design	Sample Size	Setting	Ams	Intervention Content	Primary Study Outcome	Duration
Improving BP in Colorado ¹⁴	Magid (2011)	Randomized controlled trial	338 patients randomized; 283 completed the study (138 intervention; 145 usual care) usual care)	Three healthcare systems in Denver, Colorado, including a larger health maintenance organization, a VA medical center, and a county hospital	The multimodal intervention had 4 main components: patient education, home BP monitoring, home BP measurement reporting to an interactive voice response phone system, and clinical pharmacist management of hypertension with physician oversight	3 or 4 weekly patient home-based BP monitoring; clinical pharmacist review home BP values; medication management under preapproved drug therapy management protocols	Proportion of patients who achieved guideline- recommended BP goals; change in systolic and diastolic BP between enrollment and 6- month follow-up visit	6 months
N/A ²⁴	Rifkin (2013)	Randomized, controlled effectiveness trial	43 patients (28 intervention; 15 usual care)	Chronic kidney disease and hypertension clinic at VA San Diego	Patients were randomized to use a novel telemonitoring a device pairing a Bluetooth- enabled BP cuff with an Internet- enabled hub or to usual care	Patients were instructed to measure BP at home using study devices. If a participant had consistently above-goal readings during the prior week, a study physician or pharmacist called to provide counseling or adjust medications as indicated. Additional in-person follow-up with clinic physicians or urgent care was scheduled at the discretion of the study team if it was felt that telephone counseling was not sufficient. Patients were also contacted if no	Improvement in systolic BP at 6 months	6 months

Lead Investigator Investigator Lead Investigator Investigator Pinary Su Investigator Study Title Pinary Year) Sutip Design Sample Size Setting Mary Investigator Outcome Vu-MEDIC ²⁶ Taveira (2010) Randomized 109 patients Volutiones, Florid Patients received Vu-MEDIC readings were reported Vu-MEDIC readings were reported Vu-MEDIC Settings Maria Vu-MEDIC Settings were reported	Lead Investigator Title Lead Investigator Stucy Design Stucy Design Setting A DIC ²⁵ Taveira (2010) Randomized 109 patients YA medical center in trial Frovidence, Rhode Providence, Rhode DIC ²⁵ Taveira (2010) Randomized 109 patients Providence, Rhode Providence, Rhode Vivian (2002) Randomized 56 patients (27 intervention; 29 usual VA medical center in trial T						
DIC ⁵⁵ Taveira (2010) Randomized 109 patients controlled completed the study Providence, Rhode either: 4 weekly a 40-to 60-minute reported that trial (58 intervention; 51 Island essions of the educational usual care) intervention in nutritionist, physical addition to usual therapist, or component by nues, intervention in nutritionist, physical addition to usual therapist, or care or usual therapist, or controlled intervention; 29 usual Philadelphia, were scheduled to trages inprovided to the origon or usual tradecare trade	EDIC ²⁵ Taveira (2010) Randomized 109 patients VA medical center in F controlled completed the study Providence, Rhode trial (58 intervention; 51 Island usual care) island vivian (2002) Randomized 56 patients (27 VA medical center in T controlled intervention; 29 usual Philadelphia, trial care) Pennsylvania		Setting	Ams	Intervention Content	Primary Study Outcome	Duration
Care alone by pharmacist-led behavioral and controlled intervention; 29 usual Philadelphia, am participants made appropriate trial care) Pennsylvania were scheduled changes in prescribed with a clinical dosages, and provided pharmacist. I comest in a cereiving usual effects, recommended care received lifetyle changes, and provided care received lifetyle changes, atom to filetyle changes, a	Vivian (2002) Randomized 56 patients (27 VA medical center in T controlled intervention; 29 usual Philadelphia, trial care) Pennsylvania	-	VA medical center in Providence, Rhode Island	Patients received either: 4 weekly sessions of the VA-MEDIC intervention in addition to usual care or usual	readings were reported VA-MEDIC consisted of a 40- to 60-minute educational component by nurse, nutritionist, physical therapist, or pharmacist followed	Attainment of target goals in hemoglobin A ₁ c, BP (systolic BP <130 mm Hg, diastolic BP <80 mm Hg), fasting lipids, and tobacco	1 month
from their usual compliance).	from their us	47	VA medical center in Philadelphia, Pennsylvania	The intervention arm participants were scheduled to meet monthly with a clinical pharmacist. Patients receiving usual care received standard care from their usual	by pnarmactsr-led behavioral and pharmacologic interventions The clinical pharmacist made appropriate changes in prescribed drugs, adjusted dosages, and provided drug counseling (eg, discussion of side effects, recommended lifestyle changes, assessment of compliance).	use Changes in compliance, BP, and patient satisfaction	6 months

The Adherence and Intensification of Medications (AIM) study was a cluster-randomized controlled effectiveness study that also used a pharmacist interventionist.⁶ During a 14-month period at five facilities, including two VA hospitals, primary care teams were randomized to either: (1) a program led by a clinical pharmacist trained in motivational interview-based behavioral approaches and authorized to make BP medication changes or (2) usual care. The mean systolic BP decrease from 6 months before to 6 months after the intervention period was 9 mm Hg in both arms. Mean systolic BP of eligible intervention patients were 2.4 mm Hg lower (95% CI, -3.4 to -1.5; P<.001) immediately after the intervention than those achieved by control patients.⁷

To evaluate the effectiveness of group medical clinics in the management of comorbid diabetes and hypertension, Edelman and colleagues²³ conducted a two-site randomized controlled trial. The group medical visits were comprised of seven to eight patients and a multidisciplinary care team. Groups met every 2 months. At each visit, BP was checked and home blood glucose values were collated. Patients attended an educational session delivered by the nurse or educator. Topics of the education sessions were tailored to members' needs. The pharmacist and the primary care internist reviewed patient medical records, BPs, and home blood glucose readings during each session and developed individualized plans for medication or lifestyle management directed toward improving BP and HbA_{1c} level. Mean systolic BP improved by 13.7 mm Hg in the intervention group and 6.4 mm Hg in the usual care group $(P=.011)^2$

The Veterans Affairs Multi-disciplinary Education and Diabetes Intervention for Cardiac Risk Reduction (VA-MEDIC) study evaluated whether a pharmacistled group medical visit program could improve achievement of target goals in hypertension and other goals among patients with type 2 diabetes compared with usual care.²⁵ The program involved small-group training sessions of four to eight participants lasting 40 to 60 minutes. A nurse, nutritionist, physical therapist, or pharmacist delivered the educational component, followed by pharmacist-led behavioral and pharmacologic interventions, during 4 weekly sessions. After 4 months, a greater proportion of the intervention group participants were guideline adherent for BP levels compared with the usual care arm. A greater proportion of VA-MEDIC participants vs controls achieved a systolic BP <130 mm Hg.²⁵

The extended VA MEDIC-E program²⁰ involved diabetic patients randomly assigned to standard primary care or an intervention consisting of four weekly group sessions, followed by five monthly booster group sessions. Half of the session involved diabetes-specific lifestyle education and half pharmacotherapeutic interventions performed by a clinical pharmacist. Educational topics, method of delivery, and teacher varied depending on the session. The booster intervention

focused on group needs such as discussing options for exercising during inclement weather. After 6 months, the intervention group achieved target goals for HbA_{1c} values (40.8% in cases vs 20.4% in usual care, P=.028) and systolic BP <130 mm Hg (58% cases vs 32.7% in usual care, P=.015).²⁰

Behavioral Only

Brauer²² conducted a 6-month study using relaxation therapy for improved BP (Table II). The three arms included: (1) therapist-conducted, face-to-face progressive, and deep-muscle relaxation training; (2) progressive deep-muscle relaxation therapy conducted mainly by home use of audiocassettes; or (3) nonspecific individual psychotherapy. After 6 months of postintervention follow-up, the therapist-conducted relaxation therapy group showed the greatest changes (-17.8 mm Hg systolic, -9.7 mm Hg diastolic).²²

Bosley and Allen²¹ examined the use of coping skillbuilding to reduce BP levels. Patients were randomized to receive either: (1) cognitive self-management training (CSM); (2) attention placebo control; or (3) current clinic conditions control. Participants in the CSM and attention placebo groups met for eight weekly 45minute sessions in groups of three to six members. In the CSM group, participants were trained to monitor their behavior in stressful experiences, to become aware of their bodies' reactions, and the negative self-talk that occurred in such situations. Participants in the third group received only regular clinic care. The authors found a positive association between training and reduction of systolic BP.²¹

The Veterans' Study to Improve the Control of Hypertension (V-STITCH) trial was a 2-year health services intervention.⁴ Primary care providers at one North Carolina-based clinic (n=30) were randomized to an intervention or control group. Intervention providers received a patient-specific electronically generated hypertension decision support system delivering guideline-based recommendations at each visit. Patients whose providers were in this group were randomly assigned to receive a telephone-administered intervention or usual care. The patient-level intervention involved needs assessment, followed by tailored behavioral and education models to promote medication adherence and improve specific health behaviors.⁴ Rates of BP control for all patients receiving the patient behavioral intervention improved from 40.1% to 54.4% at 24 months (P=.03); patients in the nonbehavioral intervention group improved from 38.2% to 43.9% (*P*=.38).

Roumie and colleagues^{8,9} conducted a cluster-randomized trial to examine effectiveness of three quality improvement interventions of increasing intensity in improving veterans' BP control. Randomization occurred at the provider level; providers were randomized to one of three groups: (1) education, (2) education and alert, and (3) education and alert plus patient education. Educational information was delivered to

TABLE II. Sun	nmary of Publish	ed Studies Desc	ribing VA Hyperte	nsion Control I	Interventions That	Summary of Published Studies Describing VA Hypertension Control Interventions That Use Behavioral Components	onents	
Study Title	Lead Investigator (Primary Year)	Study Design	Sample Size	Setting	Ams	Intervention Content	Primary Study Outcome	Duration
N/A ²¹	Bosley (1989)	3-arm randomized controlled trial	41 patients (14 intervention; 27 control)	VA High BP Program	Three arms: Self- Management Training (CSM), Attention Placebo Control, and Current Clinic Conditions Control	In the CSM group, the trainer presented information regarding the dynamics of stress and its role as a contributing factor in hypertension. Participants were trained to monitor their behavior in stressful experiences and be aware of their stress.	Coping style, psychological distress, and BP	1
Veterans Study to Improve the Control of Hypertension (V-STITCH) ^{3-5,10-12}	Bosworth (2005)	Randomized controlled health services intervention trial with split-plot design	30 primary care providers 588 patients (294 intervention; 294 control)	Durtham VA Medical Center Primary Care Clinic	Patient intervention included telephone contact by a nurse case manager every 2 months for 24 months	Teleboor Teleboor delivered information in nine educational and behavioral modules (for patients); audit and feedback of the provider's panel of patients with regards to guideline- recommended BP targets and medication choices, and recommendations about mangement of patients' hypertension (for providers)	Proportion of patients who achieve a BP ≤140/90 mm Hg at each outpatient clinic visit over 24 months	2 years
N/A ²²	Brauer (1979)	Three-arm single- blind randomized controlled study	29 patients (10 therapist-conducted; 9 tape-recorded; 10 control)	Palo Afto VA Hospital Outpatient Medical Clinic	Three arms: therapist- conducted, face-to- face progressive, deep-muscle relaxation training for 10 weekly sessions; progressive deep- muscle relaxation therapy conducted mainly by home use of audiocassettes; nonspecific individual psychotherapy	Therapists then instructed patients thoroughly in relaxation techniques (eg, comfortable position, systematically tensing and relaxing muscles, brancting). Patients were instructed to practice the instructed to practice the instructed to practice the rechnique at home for 20 minutes daily, using a record of their home practice, showing their progress and success in relaxing, particularly during identified stressful situations	Improvement in systolic and diastolic BP	10 weeks

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	Lead Investigator (Primary Year)	r Study Design	Sample Size	Setting	Arms	Intervention Content	Primary Study Outcome	Duration
	Mosack (2012)	Cluster randomized controlled trial	Randomized 113 volunteer leaders	58 Veterans Service Organizations in Southeast Wisconsin	Compared two hypertension self- management approaches: a peer- led intervention that occurred in the context of monthly post meetings and an educational seminar intervention covering similar information in stand- alone 90-minute presentations by health professionals	Weighing, use of pedometers, and self- monitoring through automated sphygmomanometers located at the posts	Improvement in systolic BP and hypertension knowledge, increased fruit/ vegetable intake, and pedometer use	12 months
	Roumie (2006)	Cluster randomized control trial	 182 providers (54 provider education only; 62 provider education and alert; 66 provider education, alert, and patient education); 1341 patients 	2 hospital- based and 8 community- based clinics in the VA Tennessee Valley Healthcare System	Providers were randomly assigned to receive either: (1) weekly educational e-mails only; (2) weekly educational e-mails plus patient- specific alerts; or (3) both education and alerts plus their patients were sent an interventional letter	Provider education requested reevaluation of a patient's antihypertensive regimen; alerts were one-time, patient-specific notifications sent by the pharmacy to the provider through the electronic health record; the patient letter advocated drug adherence, lifestyle modification, and conversations with providers	Proportion of patients with systolic BP ⊲140 mm Hg; intensification of antihypertensive medication	6 months
	Wakefield (2011)	3-arm randomized controlled efficacy trial	302 patients (102 low- intensity intervention; 33 high- intensity intervention; 107 usual care)	lowa City VA Medical Center	Patients in the high- intensity group had a device that used a branching disease management algorithm. The algorithm was programmed so that	The intervention combined close surveillance via a home tele-health device and nurse care management. Both intervention groups received care	Primary outcomes were hemoglobin A _{1c} and systolic BP; secondary outcome was adherence. Outcomes were measured at the end of the intervention	6 months

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Study Title	Lead Investigator (Primary Year)	Study Design	Sample Size	Setting	Arms	Intervention Content	Primary Study Outcome	Duration
					patients received	management from a study (6 months) and again	(6 months) and again	
					both standard	nurse	at 12 months to	
					prompts each day		assess maintenance	
					and a rotation of			
					questions and			
					educational content.			
					The low-intensity			
					group did not use the	τ.		
					algorithm			
Abbreviations: -, r	Abbreviations: -, not stated in the article; BP, blood pressur	le; BP, blood pressu	ire; N/A, not applicable; VA, Veterans Affairs.	;; VA, Veterans Affa	irs.			

providers via e-mail with a Web-based link to guidelines.²⁸ For providers in the alert-receiving arm, onetime, patient-specific, electronic alerts were sent by the pharmacy to the prescribing provider via the patient's electronic medical record during a 1-week period. For the third arm, a personalized letter was sent to patients containing educational information and recommended use of behavioral strategies to improve BP control. Patients of providers who were randomly assigned to the patient education group had better BP control (138/ 75 mm Hg) compared with those in the provider education and alert or provider education alone groups (146/76 mm Hg and 145/78 mm Hg, respectively).⁸

Wakefield²⁷ conducted a single-site, randomized controlled trial evaluating the efficacy of a nursemanagement home tele-health and remote monitoring intervention to improve outcomes among veteran patients with diabetes and hypertension. There were three arms: high-intensity, low-intensity, and usual care. Both intervention groups received care management from a study nurse that entailed weekday monitoring. Based on responses from patients in the intervention group, the nurse delivered follow-up in the form of providing additional health information, increased monitoring, or contacting a physician as needed. Both intervention groups were instructed to measure BP daily and blood glucose as directed by their physician. For the high-intensity group, the multidisciplinary study team developed a branching disease management algorithm that was programmed into study devices and focused on behavior modification and lifestyle adjustments. Patients in the low-intensity group were asked questions daily, but were not exposed to the branching algorithm. The high-intensity patients had a significant decrease in systolic BP compared with the other groups at 6 months and this pattern was maintained at 12 months.²⁷

The final study using behavioral strategies without medication management is unique in two ways. First, the Posts Working for Veterans Health (POWER) study^{15,16,29} was conducted in a Veterans Service Organization (VSO). In addition, rather than relying exclusively on medical professionals to serve as interventionists it uses peer leaders. All posts received a digital bathroom scale, pedometers, and automated BP monitors. The professionally led groups held three 90minutes sessions, which were advertised and repeated six times around the study area (ie, southeastern Wisconsin) so that at least one meeting was convenient for all participants. Peer leaders were trained prior to leading sessions. Their sessions were held monthly at the post and included approximately 12-minute "health corner" presentations on a specific topic such as physical activity, medication adherence, or other important topic. Peer leaders underwent eight mini-training sessions to equip them with a script to deliver the intervention. Hypertensive peer leaders lowered their systolic BP by 3.93 mm Hg (P=.04) and engaged in healthier behaviors compared with leaders from other groups.15

DISCUSSION

Several conclusions can be drawn. First, while many interventions reported in this study effectively improved BP control, they did so using a myriad of approaches. Peers, nurses, primary care providers, and pharmacists delivered interventions. Settings varied from home-based with telephone support to clinic- or community-based. Educational content, contact frequency, and intervention intensity were mixed. Many of the successful interventions relied on increased frequency of contact. As an integrated healthcare system, the VA is unique in that it bridges services related to diagnosis, treatment, medication management, and behavioral counseling, among others. As healthcare delivery evolves, with more organizations adopting an Accountable Care Organization model, this may become increasingly possible in traditionally nonintegrated healthcare settings. Additionally, the VA emphasizes quality monitoring and improvement. While not specifically addressed in the included studies, this culture and model of care may be at least partially responsible for creating an environment where these interventions can be carried out and successfully improve hypertension control.

CONCLUSIONS

While there is no universal solution to improve hypertension, there are several common ingredients among interventions that successfully improve medication adherence and often improve BP control.³⁰

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