

METHODS PAPER

The PCORnet Blood Pressure Control Laboratory

A Platform for Surveillance and Efficient Trials

BACKGROUND: Uncontrolled blood pressure (BP) is a leading preventable cause of death that remains common in the US population despite the availability of effective medications. New technology and program innovation has high potential to improve BP but may be expensive and burdensome for patients, clinicians, health systems, and payers and may not produce desired results or reduce existing disparities in BP control.

METHODS AND RESULTS: The PCORnet Blood Pressure Control Laboratory is a platform designed to enable national surveillance and facilitate quality improvement and comparative effectiveness research. The platform uses PCORnet, the National Patient-Centered Clinical Research Network, for engagement of health systems and collection of electronic health record data, and the Eureka Research Platform for eConsent and collection of patient-reported outcomes and mHealth data from wearable devices and smartphones. Three demonstration projects are underway: BP track will conduct national surveillance of BP control and related clinical processes by measuring theory-derived pragmatic BP control metrics using electronic health record data, with a focus on tracking disparities over time; BP MAP will conduct a cluster-randomized trial comparing effectiveness of 2 versions of a BP control quality improvement program; BP Home will conduct an individual patient-level randomized trial comparing effectiveness of smartphone-linked versus standard home BP monitoring. Thus far, BP Track has collected electronic health record data from over 826 000 eligible patients with hypertension who completed ≈3.1 million ambulatory visits. Preliminary results demonstrate substantial room for improvement in BP control (<140/90 mm Hg), which was 58% overall, and in the clinical processes relevant for BP control. For example, only 12% of patients with hypertension with a high BP measurement during an ambulatory visit received an order for a new antihypertensive medication.

CONCLUSIONS: The PCORnet Blood Pressure Control Laboratory is designed to be a reusable platform for efficient surveillance and comparative effectiveness research; results from demonstration projects are forthcoming.

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Uncontrolled blood pressure (BP) is a leading preventable cause of death,¹ causing over 450 000 deaths per year in the United States.^{2,3} While effective and affordable medications are available to control BP, multiple rounds of medication adjustment and intensification are typically required, and BP control is often not achieved.^{4,5} With the 2017 American College of Cardiology/American Heart Association Hypertension Guideline defining lower BP thresholds for diagnosis, treatment, and control,⁶ the prevalence of hypertension now approaches 50% of all US adults, and millions more Americans already treated for hypertension are now considered to have uncontrolled BP.^{6,7} Achieving optimal BP control at the population-level could save thousands of lives per year.⁸

It is unclear, however, how best to improve BP control rates in the United States. Some organizations have reconfigured care delivery and achieved improvements in control,^{9–13} but many of these approaches are resource intensive and may not be feasible or sustainable in all settings, particularly in resource-poor settings such as safety net clinics.^{14–16} Home BP monitoring can be effective, but generally requires additional support to produce significant and lasting reductions in BP.^{17–19} It remains unclear what types of additional support will be both effective and sustainable across varying health-care delivery settings. Emerging technologies including smartphone apps and wearables^{20–23} could facilitate BP measurement, tracking, interpretation, patient-clinician communication, medication decision-making, and adherence and thereby improve the various healthcare processes required for BP^{24,25} control, but few studies of technology effectiveness measuring BP control outcomes have been conducted.²²

Randomized controlled trials (RCTs) are required to accurately assess and compare effectiveness of different strategies for improving BP control. RCTs, however, are difficult, time-consuming, and expensive to conduct.^{26–30} The National Heart, Lung, and Blood Institute alone spent over \$90 million in 2018 (\$1.6 billion since 1985) on RCTs relevant to BP (National Institutes of Health RePORTER query³¹ and examples^{29,30}). New methods that improve RCT efficiency could help accelerate evidence generation and translation of innovative healthcare delivery solutions into major population health benefits.

Large simple real-world pragmatic trials that leverage existing resources may help streamline evidence generation and reduce burden on both investigators and patients.^{28,32–35} These methods may be particularly useful for measuring effectiveness of BP control interventions. Unlike many phenotypes, BP is measured routinely during healthcare delivery and recorded systematically in electronic health records (EHRs). Controlling High Blood Pressure, which can be constructed using data in the EHR, is a National Quality Forum endorsed performance measure (NQF 0018³⁶) recognized by the Centers

for Medicare and Medicaid Services for use in clinical incentive programs.^{37,38} While this metric has limitations, it has direct relevance to public health; demonstrating even a small average improvement in BP control from a scalable intervention would likely translate to significant health benefits when implemented broadly.³⁹

We aim to establish a platform for conducting large, simple, patient-centered real-world RCTs designed to demonstrate and compare effectiveness of BP control interventions, and a national surveillance system for monitoring both population-wide and local improvements in BP control. Our platform leverages EHR systems now ubiquitous in the United States; PCORnet, the National Patient-Centered Clinical Research Network funded by the Patient-Centered Outcomes Research Institute (PCORI);^{40,41} and the Eureka Research Platform, a National Institutes of Health-funded digital system designed to enable direct-to-participant research including collection of patient-reported outcomes and mHealth data from wearable devices and smartphones.^{42,43} In this design and methods paper, we describe the structural features of the platform—which we call the PCORnet Blood Pressure Control Laboratory (BP Control Lab)—and the 3 projects now underway using the platform that will demonstrate its utility. We end with a description of the data assets of the network of PCORnet organizations currently participating and preliminary results from our surveillance project.

STRUCTURAL FEATURES

The BP Control Laboratory brings together a set of established resources that can be leveraged to support efficient research and surveillance. Below, we describe these modular resources, and how they are configured to support large simple real-world pragmatic RCTs, local quality improvement efforts, and national surveillance. We also describe the collaborative networks that support the BP Control Lab.

The BP Control Lab uses PCORnet specifically for access to EHR data and more broadly for access to a networked clinical research infrastructure. PCORnet was jointly envisioned by PCORI and the National Institutes of Health⁴⁴ and launched in 2013 by PCORI.^{40,44,45} PCORnet's Clinical Research Networks support curation of EHR data in a common data model⁴⁶ that allows querying of EHR data across organizations using standardized queries. Patient-level data are retained locally at each organization/network and are queried via a distributed research analysis system administered by the PCORnet Coordinating Center (Figure 1). Along with systolic and diastolic BP measurements made in the context of healthcare encounters, the PCORnet Common Data Model includes information about patient demographics, encounters, diagnoses, medications, select laboratory measurements, and other domains

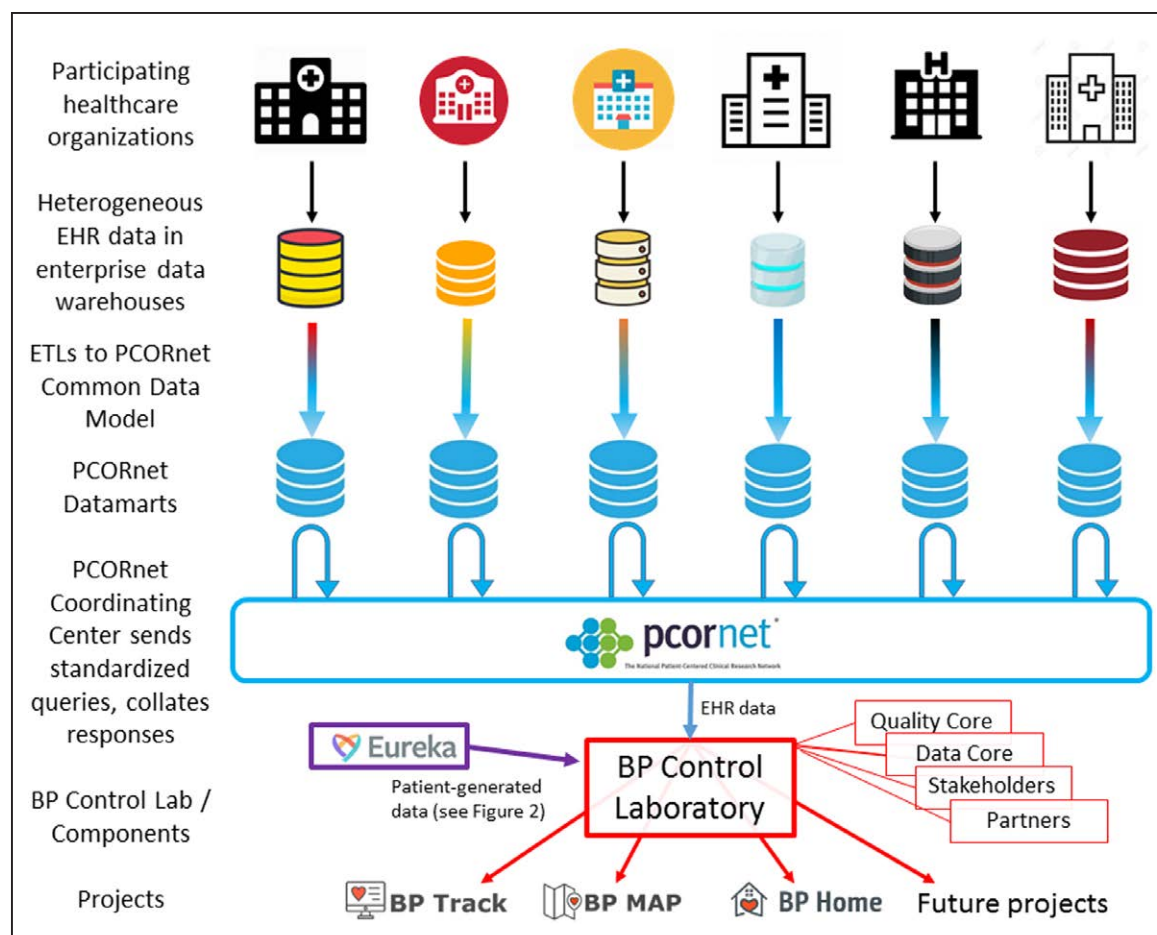


Figure 1. Electronic health record (EHR) data flow and distributed querying in PCORnet, the National Patient-Centered Clinical Research Network.

EHR data are generated by participating healthcare organizations and stored in local, heterogeneously structured comprehensive enterprise data warehouses (eg, Clarity for Epic EHR systems). To support PCORnet, each organization executes regular extract, transform, and load (ETL) operations that transform those EHR data into a homogeneous and simplified/intuitive set of relational data tables—the PCORnet Common Data Model⁴⁶—that are maintained on locally controlled servers—PCORnet Datamarts—at each participating healthcare organization (blue database icons). The data are not stored centrally by PCORnet or by the Blood Pressure (BP) Control Laboratory. To access these data, the BP Control Laboratory Data Core, with input from the quality core, partners, and stakeholders, develops common data model queries (written in SAS) that are distributed by the PCORnet Coordinating Center to each participating datamart. Datamarts then run the query locally and return results to the PCORnet Coordinating Center, which collates results and delivers them to the BP Control Laboratory. The BP Control Laboratory then links the EHR data with patient-generated data from Eureka as needed (eg, for BP Home, see Figure 2) and delivers it to the project teams supported by the Laboratory. Ongoing projects include BP Track, BP MAP, and BP Home (see Figure 3).

potentially useful in evaluating effectiveness of BP control interventions⁴⁶ and has demonstrated utility for hypertension surveillance.⁴⁷

To complement data collected during healthcare encounters, the BP Control Lab uses the Eureka Research Platform for direct patient engagement and collection of patient-generated health data (Figure 2). Eureka (originally named the Health ePeople Resource for Mobilized Research) was funded by the National Institutes of Health in 2015^{42,48} to accelerate use of mHealth data in research and evaluation of mHealth technology for improving health. Its multitenant cloud-based platform currently supports development of web- and mobile app-based patient portals that directly engage patients in eConsent, eligibility assessment, online surveys, and data collection from wearable devices and smartphones for prospective research studies including RCTs.⁴³ The platform supports secure

tracking and data linkage of Eureka enrollees recruited across different systems (eg, patients recruited from a healthcare delivery organization), retrieval of BP measurements from home BP monitoring devices via electronic data transfer, and a study management portal with customizable reports.

The BP Control Laboratory represents a collaborative effort and partnership between PCORnet entities, the American Medical Association (AMA), and the American Heart Association. The project was conceived by investigators and patients participating in the PCORnet Cardiovascular Health Collaborative Research Group,⁴⁹ including representatives from 4 Clinical Data Research Networks (OneFlorida,⁵⁰ REACHnet,⁵¹ ADVANCE,⁵² and STAR⁵³), a Patient-Powered Research Network focused on cardiovascular health (the Heart Research Alliance,⁵⁴ formerly named the Health eHeart Alliance⁵⁵), an active Patient Advisory Board, and the PCORnet Coordinating

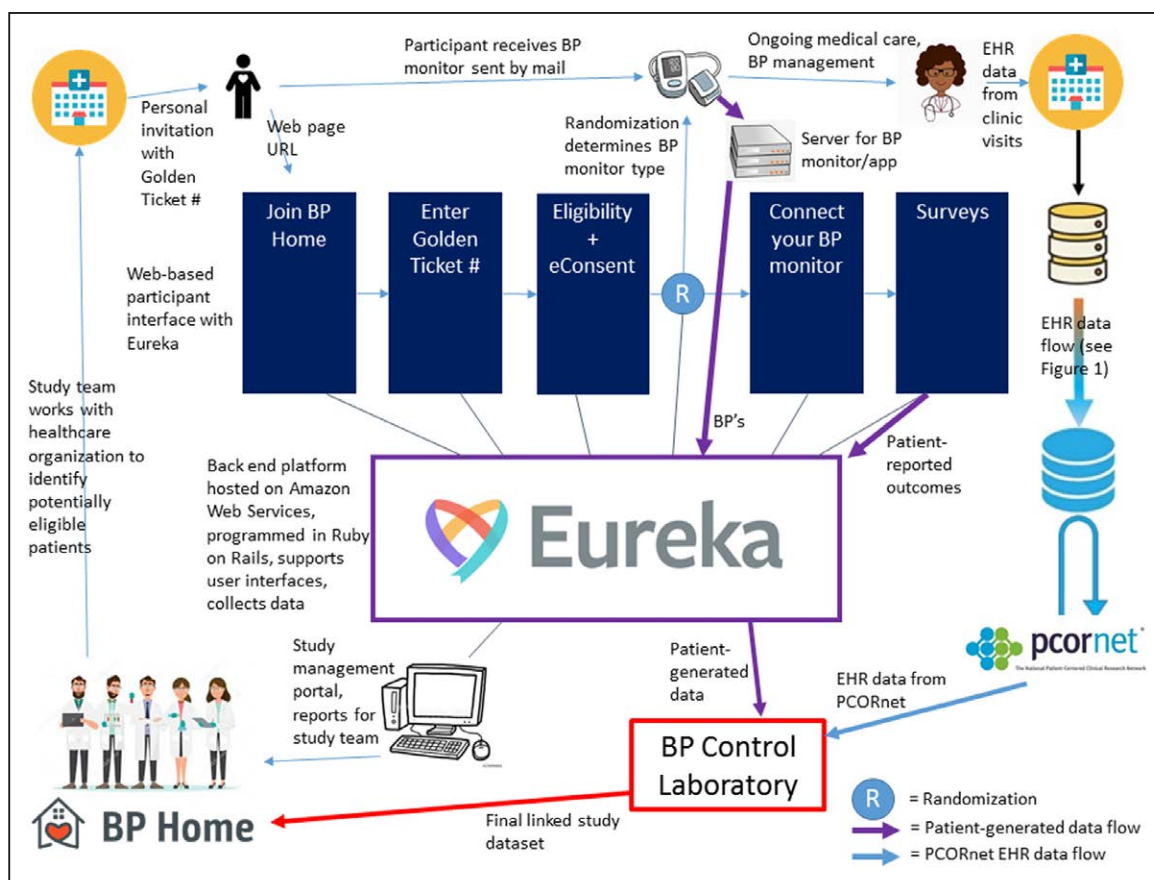


Figure 2. Patient-generated health data collection via the Eureka Research Platform.

For BP Home and future projects like it, patient-generated health data including patient-reported outcomes and blood pressure (BP) measurements from home BP monitors are collected via the Eureka Research Platform. Study teams (lower left corner) work with site personnel at healthcare organizations to identify potentially eligible patients and send them personal invitations to join the study. Interested patients go to a Eureka-hosted web portal (optimized for viewing on either a desktop or smartphone) that takes them through study information, eligibility, and eConsent procedures. They also enter a Golden Ticket number provided with the personal invitation that enables future identity linkage. Eligible and consenting participants are randomized to 1 of the 2 study arms and are mailed a home BP monitor (smartphone-linked or standard depending on randomization arm, see Figure 3), which they use for ongoing medical care and BP management with their clinician. Participants in the smartphone-linked arm then authorize connecting their home BP monitor to Eureka, allowing Eureka to obtain their BP measurements from the device company server. Participants in both arms fill out surveys that allow Eureka to gather patient-reported outcomes and other information. The study team can access reports and participant information through a study management portal hosted by Eureka. The BP Control Laboratory uses the Golden Ticket number to link patient-generated data from Eureka with electronic health record (EHR) data from Patient-Centered Outcomes Research Network (PCORnet; see Figure 1) and provides the final linked study dataset to the study team for analysis.

Center.⁴¹ To build the BP Control Laboratory, a collaborative partnership including these PCORnet entities, the AMA, and the American Heart Association applied for and received funding through PCORI's Partnerships to Conduct Clinical Research within PCORnet Funding Announcement.⁵⁶ The 3 projects funded by our award, described below, demonstrate BP Control Laboratory functionality and utility.

DEMONSTRATION PROJECTS

Overview

Our PCORI award supports 3 projects, each designed to answer scientific research questions and demonstrate different aspects of BP Control Laboratory functionality. These 3 projects, named BP Track, BP MAP, and BP Home, are described below, and summarized in Figure 3.^{57,58}

BP Track: A National Surveillance System

BP Track aims to establish a national BP control surveillance system that generates statistics on BP control and BP-related quality metrics for participating healthcare organizations. Organizations must contribute data to a PCORnet datamart that agrees to respond to quarterly queries written in SAS (Statistical Analysis Systems, Cary, NC) against the PCORnet Common Data Model, and in return receives access to metric performance reports. The queries will produce a set of quality metrics relevant to improving BP control (BP Control Metrics), including Controlling High Blood Pressure (NQF 0018^{36,59,60}) and Improvement in Blood Pressure (CMS65v7⁶¹) and additional process measures relevant to clinical management and treatment practices for BP control (Table 1). Metric design and development is guided by 2 frameworks—the AMA's MAP framework (Measure Accurately, Act

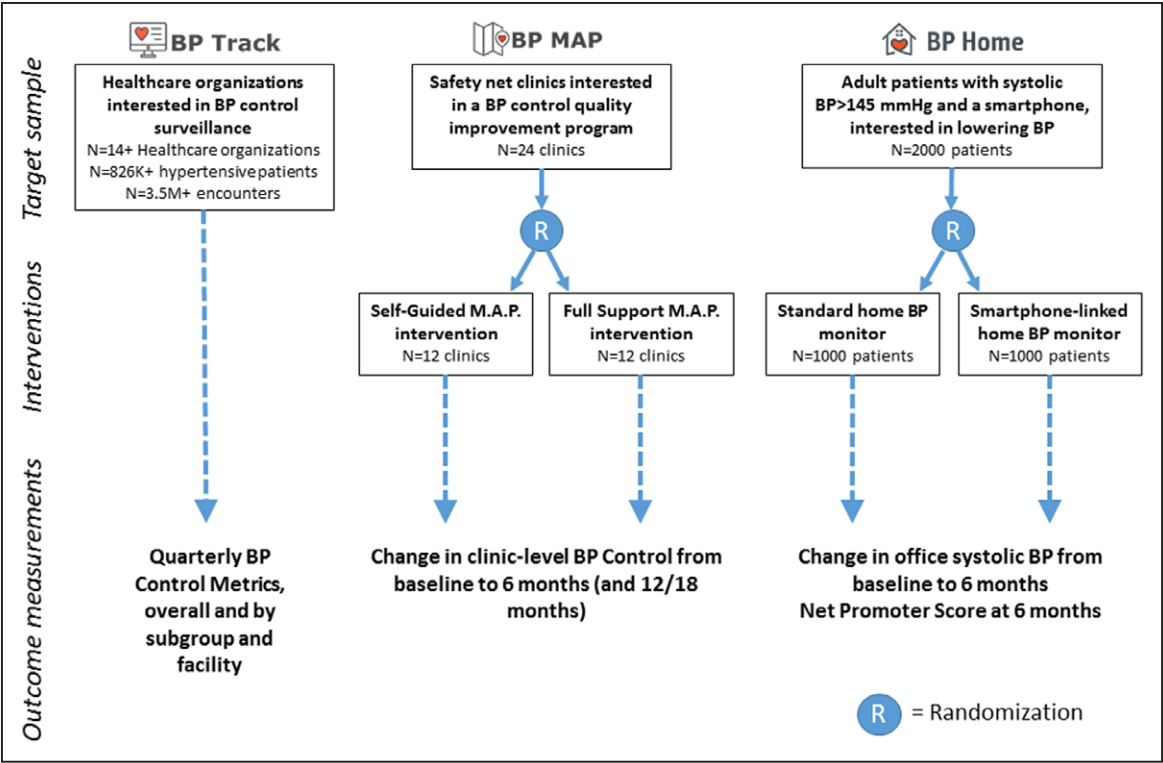


Figure 3. Design features of three projects currently supported by the Blood Pressure Control Laboratory.

The Blood Pressure (BP) Control Laboratory currently supports BP Track (a national surveillance project), BP MAP (a cluster randomized controlled trial [RCT]), and BP Home (an individual-level RCT). In the figure, we describe the target sample (target numbers and characteristics of the sample units), interventions (for the 2 RCTs), and the primary outcome measurement(s) for each project. BP control metrics, including overall BP control and 9 other BP-related quality metrics, are described in Table 1. MAP indicates Measure Accurately, Act Rapidly, Partner with Patients (a quality improvement program for BP control managed by the American Medical Association).^{24,57,58}

Rapidly, and Partner with Patients)^{24,57,58} and the Blood Pressure Control Model^{25,62}—that specify relevant clinical processes. Additional metrics perceived to be useful to stakeholders can be added over time.

BP Control Metric results will be produced overall for each participating PCORnet datamart, and for any number of individual clinical units within the datamart (eg, a particular general internal medicine clinic). Each PCORnet datamart will specify clinical units with at least one identified clinician stakeholder, to whom clinical unit-specific metrics will be provided. Each metric will be produced for the overall relevant patient population (in the datamart, or the clinical unit) and for subgroups of those patients defined by categories of age (18–44, 45–64, and 65+ years), sex (male, female, and other), and race/ethnicity (Non-Hispanic Asian, black, white, and other, and Hispanic any race). Reports and interactive data visualization will allow stakeholders to view their results and compare to blinded results from other participating organizations.

BP Track will support quality improvement efforts by providing systematic measurements of the specified quality metrics over time. Using these quality metrics, quality improvement programs can target particular processes in need of improvement, implement interventions, and use BP Track to assess change in metrics over time in relevant clinical units and patient sub-

groups compared to control units without the same exposure. This approach will be used for BP MAP (see below). BP Track will also support participation in the Target:BP Program,⁶³ a national initiative formed by the American Heart Association and the AMA that aims to help healthcare organizations prioritize and improve BP control and recognize organizations for achieving BP control rates of 70% or higher.

Although organizations participating in BP Track will not represent a random sample of either the US population or US healthcare organizations, the relatively broad participation in the program (currently 14 PCORnet datamarts with healthcare organizations in 15 different states) and scalability (additional organizations can participate if they support the PCORnet Common Data Model and participate in the distributed research network) makes it a potentially useful platform for national surveillance. BP Track represents the first use of PCORnet for national surveillance, and a testbed for PCORnet surveillance methodology.

BP MAP: a Cluster Randomized Quality Improvement Trial

BP MAP (Improving Blood Pressure Control in Diverse Populations by Measuring Accurately, Acting Rapidly,

Table 1. BP Control Metrics to be Tracked by the PCORnet BP Control Laboratory

No.	Metric	Description	Implementation Plan
1	BP control, % of patients	This overall measure of BP control implements NQF 0018, which defines BP control as the percent of eligible patients with hypertension for whom the BP measurements at their most recent ambulatory care visit were at goal, defined as SBP<140 mm Hg and DBP<90 mm Hg.	Wave 1
2	BP control to 2017 guideline goal, % of patients	This alternative overall measure of BP control is identical to Metric 1, except that attainment of BP control is defined by SBP<130 mm Hg and DBP<80 mm Hg, as per the goal stated in the 2017 American College of Cardiology/American Heart Association Hypertension Guideline. Note that while the treatment threshold varies in the guideline, depending on cardiovascular risk, the goal applies to all patients.	Wave 2
3	Improvement in BP, % of patients	This overall measure of BP improvement implements CMS065v4, which defines BP improvement as either a reduction of 10 mmHg in SBP or achievement of SBP that is adequately controlled (SBP<140 mmHg) in months 10 to 12 of the measurement period, among patients with hypertension not previously controlled.	Wave 1
4	Confirmatory repeated BP measurement, % of visits	This process measure is designed to capture the practice of repeating a BP measurement in the same visit when the first measurement done in clinic is high (SBP≥140 mmHg or DBP≥90 mmHg).	Wave 1
5	Terminal digit=zero, % of measurements	Inappropriate rounding of BP measurements (usually to zero) leads to measurement error and worse treatment decisions. This metric is designed to measure the extent of this behavior, which would lead to a terminal digit of zero >10% of the time (if an automated BP monitor is used) or >20% (if a manual BP monitor is used with recommended rounding to even digits). Unlike most of our metrics, lower is better, down to an ideal value of 10%–20%, which would be expected if no rounding were occurring.	Wave 2
6	Medication intensification, % of visits	This process measure captures the proportion of visits where BP is uncontrolled where a BP medication is ordered that is of a different class of medication than had previously been used. Note that this explicitly does not give credit for ordering a simple refill or medication dose increase or use of a different medication in the same class.	Wave 1
7	Repeat visit in 4 wks after uncontrolled HTN, % of visits	This process measure captures the proportion of visits by persons with uncontrolled hypertension that were followed by a subsequent visit within 4 weeks.	Wave 2
8	Average SBP reduction after medication intensification, mm Hg	This continuous metric describes the change in SBP observed between a visit with a medication intensification to the subsequent visit occurring at least 10 days later. We will collect both the average and the SD for this metric.	Wave 1
9	Use of a CCB or thiazide or thiazide-like diuretic among African-American patients on at least one medication, % of patients	Use of CCBs or a thiazide or thiazide-like diuretic medication class is recommended to treat black or African-American patients as first line monotherapy due to increased efficacy. This metric, which is limited to African-American patients with a diagnosis of hypertension taking at least one medication class, describes the prevalence of those receiving the recommended drug class.	Wave 2
10	Use of fixed dose combination product among patients taking 2 or more classes of medications, % of patients	Use of fixed dose combination medications helps with adherence, promotes rational combinations of medications, and increases likelihood of achieving BP control. This metric, which is limited to patients taking more than one BP medication class, describes the prevalence of fixed dose combination pill use.	Wave 2

BP indicates blood pressure; CCB, calcium channel blocker; DBP, diastolic BP; PCORnet, National Patient-Centered Outcomes Research Network; and SBP, systolic BP.

and Partnering with Patients) is a cluster randomized RCT that will compare effectiveness of a Full Support versus a Self-Guided version of a clinic-level hypertension quality improvement intervention. The quality improvement intervention is based on the AMA's M.A.P. framework²⁴ and 6-month M.A.P. BP Improvement Program, which includes quality improvement materials and a protocolized program with support from dedicated practice change facilitators. Interventions based on the M.A.P. framework have shown evidence of effectiveness, with improved BP control and process metrics in prepost analyses^{57,58} with sustainability at 12 months.⁵⁷ It is unclear, however, how much the support

from the dedicated practice change facilitators (who are trained centrally by the AMA in a train-the-trainer model) is required for the program to be successful.

BP MAP will randomize 24 clinics from 2 PCORnet research networks (REACHnet and OCHIN) to 1 of the 2 versions of the AMA's M.A.P. BP Improvement Program: Full Support versus Self-Guided. The Full Support arm will include an on-site practice assessment, an in-person launch meeting, training, and personalized support for dedicated practice change facilitators from experienced AMA staff and access to an online Digital Guide containing resources and training materials. The Self-Guided arm will have access to the Digital Guide and an informa-

tional webinar at launch but will not receive in person site visits or practice change facilitator support. Both arms will run BP Track queries (Table 1) on a monthly basis and use the results to guide and target their quality improvement efforts. The primary outcome will be change in BP control (NQF 0018) from baseline to 6 months, as assessed using BP Track queries of the PCORnet Common Data Model (Table 1, Metric 1). This outcome will be compared between randomized arms; and each arm will also be compared against Usual Care (nonrandomized), which will include other PCORnet datamarts participating concurrently in BP Track (excluding datamarts with clinical units participating in BP MAP). Additional features of BP MAP are published on www.clinicaltrials.gov.⁶⁴ The protocol, including a description of the Digital Guide and resources required to implement the Full Support versus the Self-Guided versions of the intervention at scale, will be published separately.

BP Home: a Randomized Controlled Trial of Home Monitoring Technology

BP Home (The PCORnet Blood Pressure Home Monitoring Study) will compare effectiveness of smartphone-linked versus standard home BP monitors for helping patients with uncontrolled hypertension achieve a reduction in systolic BP. Home BP monitoring by itself has been shown to have only a small overall impact on BP control^{17,19}; home monitoring combined with additional support seems to provide more robust gains in BP control.^{17–19} While additional support that requires reconfiguration of care and extra resources may not be achievable in low-resource settings, emerging technol-

ogy that is more user-friendly in diverse patient populations (including elderly and low-income patients who tend to be late adopters of technology) may help bridge this gap.^{65–67} Developing and testing new devices, smartphone apps, and other support systems that effectively engage patients in home BP monitoring and help patients and clinicians achieve subsequent BP control remains an active area of research and development in the public and private sectors; while consumer-focused technology reviews are available,⁶⁸ these typically do not rely on evidence produced by robust comparative effectiveness methods. It is critically important to demonstrate and compare effectiveness of emerging technologies designed to enhance BP control.

BP Home will test whether having a smartphone-linked home BP monitor (that connects via Bluetooth to a smartphone and works in tandem with a commercially available smartphone app) constitutes additional support that improves BP control more effectively than a standard device without Bluetooth connectivity. Our pragmatic design features online enrollment, eConsent, and survey delivery via the Eureka Research Platform (Figure 2), simple eligibility criteria (systolic BP >145, stated desire to lower systolic BP by 10 mm Hg, and owning a smartphone), scalable interventions (2 consumer devices) provided with minimal study-specific support, and an imperfect but pragmatic outcome (reduction in systolic BP measured at most recent clinic visit, at 6 months after enrollment). Four PCORnet networks will help recruit the planned 2000 BP Home participants required for the study and will query their PCORnet datamarts for office-based BP measurements from EHR data, which will be linked with patient-report-

Table 2. Total Patients and Encounters Available for Analysis in Wave 1 of BP Track

Patient/Encounter Type	Total Numbers of Patients and Observations From Participating Datamarts	
	Median (Range) of Totals Within Each Datamart (N=14 Datamarts)	Total Across All Datamarts
Patients		
All adult patients with hypertension meeting eligibility criteria* for BP control metrics	35 719 (1042–178 132)	826 392
With diabetes mellitus diagnosis†	7826 (206–63 327)	240 753
With coronary heart disease diagnosis†	5261 (43–29 116)	112 456
With heart failure diagnosis†	2187 (24–12 993)	47 677
With depression diagnosis†	5484 (260–31 717)	116 626
With COPD diagnosis†	2067 (51–12 008)	50 788
Encounters		
All ambulatory encounters made by eligible patients*	130 956 (3704–757 235)	3 570 311
With a BP measurement available	119 078 (3655–714 894)	3 103 423

BP Track represents PCORnet Blood Pressure Control Registry. BP indicates blood pressure; COPD, chronic obstructive pulmonary disease; and PCORnet, National Patient-Centered Outcomes Research Network.

*During a defined 1-year measurement period, the following criteria are met: at least 1 ambulatory visit occurs, patient is age 18–85 y at the end of the period; a diagnosis of hypertension during the first 6 mo of the period or at any time prior; no hospice services provided to the patient; no diagnosis of end-stage renal disease, dialysis, or renal transplant during or prior; no diagnosis of pregnancy; not residing in a long-term care facility.³⁶

†Defined by a diagnosis assigned during the 1-year measurement period.

ed outcomes and home BP measurements collected by Eureka (Figure 2). The Patient Advisory Board helped design the protocol and the web portal hosted on the Eureka platform, reviews all patient-facing material, and advises the study team on all matters impacting participant experience. Additional features of BP Home are published on www.clinicaltrials.gov.⁶⁹

Description of BP Control Laboratory Data Assets and Preliminary Results From BP Track, Wave 1

Fourteen Wave 1 datamarts fully executed contracts with the PCORnet Coordinating Center in time to participate in the first quarterly BP Track queries; additional datamarts are expected to participate in subsequent Waves, which are currently funded to continue through mid-2021. Table 2 describes the totals and ranges in the number of observations (eligible patients and ambulatory visits) contributed by each Wave 1 datamart. A total of 826 392 adult patients with hypertension met eligibility criteria across datamarts, many with comorbid diagnoses relevant to hypertension control and cardiovascular disease. Those patients completed over 3 million qualifying ambulatory visits where BP was measured during the year-long observation periods selected by each datamart. One datamart reported results from 4 specific clinical units of interest within their datamarts; more will participate in this aspect of BP Track in Wave 2.

BP Control Metrics from datamarts reporting in Wave 1 showed substantial room for improvement in BP control and relevant clinical processes (Table 3). Overall results (weighted for numbers of observations) demon-

strated that 58% of adult patients with hypertension were controlled to <140/90 mmHg, confirmatory re-measurement of BP after a high ambulatory measurement was uncommon (16%), and only 12% of patients with a high BP at an ambulatory visit were subsequently prescribed a new class of BP medication. When this occurred, however, the average reduction in systolic BP was relatively large (14.6 mmHg). We observed substantial variation in these metrics between datamarts, and between clinical units within datamarts, which indicates marked room for improvement. BP Track data will be available for analysis and publication, subject to a forthcoming BP Track Publication Policy; requests may be submitted to the corresponding author.

SUMMARY

The BP Control Laboratory represents a new national infrastructure for BP control surveillance, evaluation of healthcare quality improvement efforts, and pragmatic RCTs. Results from the 3 demonstration projects will be reported in coming years. Along with the main scientific outcomes, each project will include assessments of engagement from target stakeholders including patients and trial participants, clinicians, local policy-makers (eg, medical system leadership), and researchers interested in BP control. We also plan to analyze efficiency and sustainability of the BP Control Laboratory and potential cost savings to research projects using the core infrastructure (PCORnet and Eureka). Our goal is to provide results useful to our stakeholders, and to reuse the BP Control Laboratory infrastructure that now exists for future efficient research that helps improve BP

Table 3. Aggregate Blood Pressure Control Metrics in BP Track, Wave 1

BP Control Metric		Result			
No.	Name	N*	Weighted Result*	Datamart Range† (Min–Max) N=14 Datamarts; n=826 392 Patients	Clinical Unit Range‡ (Min–Max) N=4 Clinical Units‡ n=144 432 Patients
1	Blood pressure control, % of patients	826 392	58%	40%–65%	31%–60%
3	Improvement in blood pressure, % of patients	213 240	28%	16%–37%	13%–28%
4	Confirmatory repeated blood pressure measurement, % of visits	254 820	16%	0%–92%	14%–51%
6	Medication intensification, % of visits	244 526	12%	0.5%–16%	8%–11%
8	Average SBP reduction after medication intensification, mmHg	14 928	14.6 mmHg	11.1–16.5	11.2–16.1

BP indicates blood pressure; PCORnet, the National Patient-Centered Clinical Research Network; and SBP, systolic BP.

*Overall results are calculated as weighted averages of datamart-specific results, weighted by the total number of observations (patients or visits) meeting eligibility criteria for metric calculation (N). Note: For confidentiality, all counts reported to the BP control laboratory (both numerators and denominators in proportion metrics) are masked when cell sizes are between 1 and 10. Results are reported as missing when the denominator (N) is <11. For metrics 1, 3, 4, and 6, which are proportions, we imputed a numerator of 5 if the denominator was 100 or greater and the numerator was between 1 and 10. N's represent total eligible observations contributing to nonmissing (including imputed) results.

†Datamart range represents the minimum value and the maximum value across the 14 reporting PCORnet datamarts. Two values were imputed for Metric 4, as described in the preceding footnote.

‡Clinical Unit Range represents the minimum value and the maximum value across the 4 individual clinical units within PCORnet datamarts specifically tracked in BP Track by specifying FACILITYID in the ENCOUNTER table of the PCORnet Common Data Model.

control, reduce disparities, guide evidence-based use of technology, and improve cardiovascular outcomes for the US population. Investigators interested in using the BP Control Laboratory are encouraged to contact the study authors.

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Disclosures

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