1 Clinical Policy: Critical Issues Critical Issues in the Evaluation and Management of Adult Patients 2 Presenting to the Emergency Department Asymptomatic Elevated Blood Pressure 3 This DRAFT is EMBARGOED - Not for Distribution 4 5 6 From the American College of Emergency Physicians Clinical Policies Subcommittee (Writing Committee) on 7 Asymptomatic Hypertension: 8 9 Seth Gemme, MD (Writing Committee Chair) 10 Andrew C. Meltzer, MD Richard Byyny, MD, MSc (Methodologist) 11 12 Deborah B. Diercks, MD, MSc (Committee Chair) 13 14 15 Members of the American College of Emergency Physicians Clinical Policies Committee (Oversight Committee): 16 17 Deborah B. Diercks, MD, MSc (Chair 2021-2024) 18 John D. Anderson, MD 19 Richard Byyny, MD, MSc (Methodologist) 20 Christopher R. Carpenter, MD, MSc Benjamin W. Friedman, MD (Methodologist) 21 22 Seth R. Gemme, MD 23 Charles J. Gerardo, MD, MHS 24 Steven A. Godwin, MD 25 Benjamin W. Hatten, MD, MPH 26 Jason S. Haukoos, MD, MSc (Methodologist) 27 Amy Kaji, MD, MPH, PhD (Methodologist) 28 Heemun Kwok, MD, MS (Methodologist) 29 Bruce M. Lo, MD, MBA, RDMS 30 Sharon E. Mace, MD 31 Amal Mattu, MD 32 Susan B. Promes, MD, MBA 33 Kaushal H. Shah, MD 34 Richard D. Shih, MD 35 Scott M. Silvers, MD Andrea Slivinski, RN, DNP (ENA Representative 2021-2024) 36 37 Michael D. Smith, MD, MBA 38 Molly E. W. Thiessen, MD 39 John T. Thompson, MD (EMRA Representative 2023-2024) 40 Christian A. Tomaszewski, MD, MS, MBA Stacy A. Trent, MD, MPH (Methodologist) 41 42 Jonathan H. Valente, MD Lauren M. Westafer, DO, MPH, MS 43 Stephen P. Wall, MD, MSc, MAEd (Methodologist) 44 45 Yanling Yu, PhD (Washington Advocates for Patient Safety) 46 Michelle P. Lin, MD, MPH, MS (Liaison with the ACEP Quality and Patient Safety Committee and E-QUAL 47 Steering Committee) 48 John T. Finnell, MD (Board Liaison 2020-2024) 49 Travis Schulz, MLS, AHIP, Staff Liaison, Clinical Policies Committee and Writing Committee on Asymptomatic

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### **ABSTRACT**

This clinical policy from the American College of Emergency Physicians addresses key issues in the evaluation and management of adult emergency department patients presenting with asymptomatic hypertension.

A writing committee conducted a systematic review of the literature to derive evidence-based recommendations to answer the following clinical-question: In adult emergency department patients being discharged with asymptomatic elevated blood pressure, is initiation of outpatient antihypertensive medications from the emergency department safe and effective? Evidence was graded and recommendations were made based on the strength of the available data.

### INTRODUCTION

Approximately half of adults in the United States (119.9 million) are affected by hypertension, but only 25% (27.0 million) of these individuals effectively control their blood pressure. Hypertension, defined as blood pressure greater than 130/80 mm Hg, is the primary risk factor for cardiovascular disease and good blood pressure control reduces the likelihood of subsequent stroke and heart attack. There are just over 6 million emergency department (ED) visits annually in the United States for a primary chief complaint of hypertension and of those patients, about 64% receive a primary diagnosis of hypertension.

In general, ED physicians excel at identifying acute life-threatening emergencies like stroke or myocardial infarction but have less experience with the long-term treatment for chronic illness such as asymptomatic hypertension. Wide variation in practice patterns exist for the management of patients with asymptomatic elevated blood pressure in the ED, despite the reliability of blood pressure measurements taken in the ED.<sup>6,7</sup> The benefits of starting or modifying blood pressure medications in asymptomatic high blood pressure may be countered by the potential risks. For example, some ED physicians believe that blood pressure treatment should be left to the primary care provider because of the need for long-term management and titration. Other ED physicians believe that treating asymptomatic high blood pressure in the ED represents an opportunity to improve medication compliance. The 2013 ACEP Clinical Policy did not recommend for routine ED medical interventions for asymptomatic elevated blood pressure, unless the patient had poor follow-up or the patient was part of a select high-risk patient population.<sup>8</sup>

This current ACEP clinical policy updates the 2013 clinical policy by incorporating new evidence with the aim of providing guidance for ED physicians to determine if initiation of antihypertensive medications at and/or prior to discharge from the emergency department is safe and effective.

### METHODOLOGY

This ACEP clinical policy was developed by ED physicians with input from medical librarians and a patient safety advocate and is based on a systematic review and critical, descriptive analysis of the medical literature and is reported in accordance with Preferred Reporting Items for Systematic reviews and Meta-Analyses (PRISMA) guidelines.<sup>9</sup>

### Search and Study Selection

This clinical policy is based on a systematic review with critical analysis of the medical literature meeting the inclusion criteria. Searches of PubMed, SCOPUS, Embase, Web of Science, and the Cochrane Database of Systematic Reviews were performed by a second librarian. Search terms and strategies were peer reviewed by a second librarian. All searches were limited to human studies published in English. Specific key words/phrases, years used in the searches, dates of searches, and study selection are identified under the critical question. In addition, relevant articles from the bibliographies of included studies and more recent articles identified by committee members and reviewers were included.

Using Covidence (Covidence, Melbourne, Australia), 2 subcommittee members independently reviewed the identified abstracts to assess for possible inclusion. Of those identified for potential inclusion, each full-length text was reviewed for eligibility. Those identified as eligible were subsequently abstracted and forwarded to the committee's methodology group (emergency physicians with specific research methodological expertise) for methodological grading using a Class of Evidence framework (Appendix E1).

### Assessment of Risk of Bias and Determination of Classes of Evidence

Each study identified as eligible by the subcommittee was independently graded by two methodologists.

Design 1 represents the strongest possible study design to answer the critical question, which relates to whether the

focus was therapeutic, diagnostic, or prognostic, or a meta-analysis. Subsequent design types (i.e., Design 2 and Design 3) represent respectively weaker study designs. Articles are then graded on dimensions related to the study's methodological features and execution, including but not limited to randomization processes, blinding, allocation concealment, methods of data collection, outcome measures and their assessment, selection and misclassification biases, sample size, generalizability, data management, analyses, congruence of results and conclusions, and potential for conflicts of interest.

Using a predetermined process that combines the study's design, methodological quality, and applicability to the critical question, 2 methodologists independently assigned a preliminary Class of Evidence grade for each article. Articles with concordant grades from both methodologists received that grade as their final grade. Any discordance in the preliminary grades was adjudicated through discussion which involved at least 1 additional methodologist, resulting in a final Class of Evidence assignment (i.e., Class I, Class II, Class III, or Class X) (Appendix E2). Studies identified with significant methodologic limitations and/or ultimately determined to not be applicable to the critical question received a Class of Evidence grade "X" and were not used in formulating recommendations for this policy. However, content in these articles may have been used to formulate the background and to inform expert consensus in the absence of evidence. Question-specific Classes of Evidence grading may be found in the Evidentiary Table included at the end of this policy.

### Translation of Classes of Evidence to Recommendation Levels

Based on the strength of evidence for each critical question, the subcommittee drafted the recommendations and supporting text synthesizing the evidence using the following guidelines:

**Level A recommendations.** Generally accepted principles for patient care that reflect a high degree of scientific certainty (e.g., based on evidence from one or more Class of Evidence I, or multiple Class of Evidence II studies that demonstrate consistent effects or estimates).

**Level B recommendations.** Recommendations for patient care that may identify a particular strategy or range of strategies that reflect moderate scientific certainty (e.g., based on evidence from one or more Class of Evidence II studies, or multiple Class of Evidence III studies that demonstrate consistent effects or estimates).

Level C recommendations. Recommendations for patient care that are based on evidence from Class of Evidence III studies or, in the absence of adequate published literature, based on expert consensus. In instances where consensus recommendations are made, "consensus" is placed in parentheses at the end of the recommendation.

There are certain circumstances in which the recommendations stemming from a body of evidence should not be rated as highly as the individual studies on which they are based. Factors such as consistency of results, uncertainty of effect magnitude, and publication bias, among others, might lead to a downgrading of recommendations. When possible, clinically-oriented statistics (e.g., likelihood ratios [LRs], number needed to treat) are presented to help the reader better understand how the results may be applied to the individual patient. This can assist the clinician in applying the recommendations to most patients but allow adjustment when applying to patients with extremes of risk (Appendix E3).

#### Evaluation and Review of Recommendations

Once drafted, the policy was distributed for internal review (by members of the entire committee) followed by external expert review and an open comment period for all ACEP membership. Comments were received during a 60-day open comment period with notices of the comment period sent electronically to ACEP members, published in *EM Today*, posted on the ACEP Web website, and sent to other pertinent physician organizations. The responses were used to further refine and enhance this clinical policy, although responses do not imply endorsement. Clinical policies are scheduled for revision every 3 years; however, interim reviews are conducted when technology, methodology, or the practice environment changes significantly.

### Application of the Policy

This policy is not intended to be a complete manual on the evaluation and management of adult patients with asymptomatic hypertension but rather a focused examination of critical questions that have particular relevance to the current practice of emergency medicine. Potential benefits and harms of implementing recommendations are briefly summarized within each critical question.

It is the goal of the Clinical Policies Committee to provide evidence-based recommendations when the scientific literature provides sufficient quality information to inform recommendations for a critical question. In accordance with ACEP Resolution 56(21), ACEP clinical policies do not use race-based calculators in the formulation of recommendations. When the medical literature does not contain adequate empirical data to inform a critical question, the members of the Clinical Policies Committee believe that it is equally important to alert emergency physicians to this fact.

This clinical policy is not intended to represent a legal standard of care for emergency physicians. Recommendations offered in this policy are not intended to represent the only diagnostic or management options available to the emergency physician. ACEP recognizes the importance of the individual physician's judgment and patient preferences. This guideline provides clinical strategies for which medical literature exists to inform the critical questions addressed in this policy. ACEP funded this clinical policy.

Scope of Application. This guideline is intended for physicians working in the ED.

*Inclusion Criteria.* Adult patients aged 18 years or older who present to the ED with asymptomatic elevated blood pressure without signs and symptoms of acute target organ injury.

Exclusion Criteria. Patients who present to the ED with signs or symptoms of acute hypertensive emergencies (ie, patients with clinical findings that suggest acute target organ injury such as acute stroke, cardiac ischemia, pulmonary edema, encephalopathy, and congestive heart failure), pregnant patients, patients with end-stage renal insufficiency, emergent conditions that are likely to cause elevated blood pressure not directly related to acute target organ injury (eg, trauma, other pain syndromes), and acute presentations of serious medical conditions associated with hypertension such as stroke, myocardial infarction, and congestive heart failure.

### **CRITICAL QUESTIONS**

1. In adult emergency department patients being discharged with asymptomatic elevated blood pressure, is initiation of outpatient antihypertensive medications from the emergency department safe and effective?

**Patient Management Recommendations** 

Level A recommendations. None.

Level B recommendations. None.

190	Level C recommendations. Consider the initiation of outpatient antihypertensive medications for patients
191	being discharged from the ED with asymptomatic elevated blood pressure.
192	Patients with asymptomatic elevated blood pressure should be referred for outpatient follow-up
193	(Consensus recommendation).
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### Potential Benefit of Implementing the Recommendations:

- Improvement in cardiovascular and cerebrovascular risk.
- Initiation of treatment sooner.
- Potential reduction in healthcare disparities.

### Potential Harm of Implementing the Recommendations:

- Adverse effect of the medication.
- Treating of a falsely elevated blood pressure and thus creating hypotension.

<u>Key words/phrases for literature searches:</u> Antihypertensive, Antihypertensive Agent, Antihypertensive Agents, Antihypertensive Therapy, Asymptomatic, Blood Pressure, Clevidipine, Discharge, Discharge Planning, Elevated Blood Pressure, Emergency Department, Emergency Medicine, Emergency Service, Enalaprilat, Esmolol, Fenoldopam, Glyceryl Trinitrate, High Blood Pressure, Hospital Discharge, Hydralazine, Hypertension, Labetalol, Nicardipine, Nitroglycerin, Nitroprusside, Nitroprusside Sodium, Patient Discharge, Phentolamine, Pulmonary Hypertension, and variations and combinations of key words/phrases. Searches included January 2011 to the search dates of August 23 and 24, 2022 (Appendix E4).

Study Selection: One thousand seventeen articles were identified in the searches. Six hundred sixty-seven articles were selected from the search results as candidates for further review. After grading for methodological rigor, zero Class I studies, zero Class II studies, and 1 Class III study was included for this critical question (Appendix E5).

Managing a chronic condition beyond discharge from the emergency department carries potential risks due to the episodic nature of emergency medicine. Emergency physicians might hesitate to initiate chronic medications due to both limited expertise in this area and concerns about the ongoing monitoring of the medication's safety and effectiveness. Yet, considering the widespread challenges in accessing healthcare in the United States, the ED visit might represent the sole opportunity for timely intervention. There is limited high-quality evidence directly addressing the critical question.

In the only Class III study identified, Brody et al. found that prescribing antihypertensive medication upon discharge from the ED was associated with short-term lowering of blood pressure without any increase in adverse events (Table 1).<sup>10</sup> In this retrospective analysis of 2 prospective, longitudinal randomized controlled trials, uncontrolled blood pressure was defined as greater than 140/90 mm Hg or 160/90 mm Hg, depending on which of the 2 RCTs ("PCCD" NCT00689819 and "adDReaCH" NCT01360476). Patients were included if they were

asymptomatic and excluded if they had a cardiovascular or neurovascular event or history of cardiovascular disease were excluded. Antihypertensive medications were initiated by the ED provider (Table 2). There was a total of 217 patients of which 124 were female (57%). Importantly, 208 (96%) of the patients were African American, and 65 (86%) had established hypertension at the time of the ED visit. The patients that received the antihypertensive prescription from the ED had a reduction of 11 mm Hg in blood pressure at follow up (95% CI 17 to 4 mm Hg). Both groups, with and without antihypertensive prescription, had similar rates of adverse events (1.59 versus 1.43; difference=0.16, 95% CI -0.34 to 0.67). No new neurological deficits, ischemic events, life threatening anaphylactic reactions or clinically significant hypotension (SBP <100 mm Hg) were reported in either group. These studies are consistent with Joint National Committee (JNC 8) guidelines that recommend treating hypertensive persons aged more than 60 years to a blood pressure goal of less than 150/90 mm Hg based on strong evidence and, treating a blood pressure of less than 140/90 mm Hg for other groups based on expert opinion.

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## Table 1. Adverse Events Related to the Administration of Antihypertensive Therapy. 11

### **Major adverse events:**

- (1) death from coronary heart disease (CHD);
- (2) death from other cardiovascular disease (CVD) including stroke;
- (3) death from other causes;
- (4) nonfatal myocardial infarction;
- (5) nonfatal stroke;
- (6) congestive heart failure;
- (7) surgery for aortic aneurysm;
- (8) coronary artery bypass surgery;
- (9) coronary angioplasty;
- (10) thrombolytic therapy; or
- (11) hospitalization for unstable angina

### Other adverse events defined a priori as outcome variables:

- (1) hospitalization for cerebral transient ischemic attacks (TIAs);
- (2) definite angina or intermittent claudication by Rose questionnaire; and
- (3) peripheral arterial occlusive disease defined as absent or diminished pedal pulses on one side with a bruit in the femoral artery on that side or absent or diminished pulse in any artery (femoral, posterior tibial, or dorsalis pedis) with ischemic ulcers, or history of surgery for peripheral arterial insufficiency.

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## Table 2. Class of antihypertensive medications prescribed. 10

Drug Class	Prevalence in Study
Thiazide-like diuretics	54%

Angiotensin-converting enzyme inhibitors	26%
Calcium channel blockers	10%
Beta blockers	6%

### Summary

The previous ACEP Clinical Policy discouraged routine intervention in the ED, except for specific populations, following a consensus recommendation. However, a recent review of current literature revealed a study demonstrating both efficacy and safety in treating patients with elevated blood pressure initiated from the ED. Considering this study's findings, there appears to be merit in contemplating the commencement of treatment for individuals arriving at the ED with asymptomatic elevated blood pressure.

## Future Research

Given only 1 study was identified of quality, more research is needed to better answer the critical question.

Also, future research should seek to address the following:

• Are there certain patient demographics that impact the initiation of anti-hypertensive medications from the ED?

 • What are the potential barriers and facilitators that impact the initiation of blood pressure management from the ED?

Does the availability of timely outpatient follow-up impact short- or long-term efficacy and safety in prescribing from the ED?
What is the appropriate outpatient follow-up time frame after discharging from the ED?

• For those without an established diagnosis of hypertension, is initiation of outpatient antihypertensive medications from the ED safe and effective?

## Quality Measures and Aims

ACEP uses an evidence-based approach to develop quality measures targeting variations in emergency care. ACEP's approach links measures to patient outcomes, reducing clinician burden and delivering meaningful information to clinicians and patients. Working with the ACEP Quality and Patient Safety Committee and Clinical Emergency Data Registry Committee (CEDR), the Clinical Policies Committee identified and elected to include Quality Payment Program (QPP) measure: *QPP317 Preventive Care and Screening: Screening for High Blood Pressure and Follow-Up Documented* (Appendix E6). The aims of this measure are:

- 1. Increase the percentage of patients 18 years and older screened for high blood pressure during the measurement period.
- 2. Discharge the patient with a documented follow-up plan if the result of the blood pressure screening is pre-hypertensive or hypertensive.

Relevant industry relationships: There were no relevant industry relationships disclosed by the subcommittee members for this topic.

Relevant industry relationships are those relationships with companies associated with products or services that significantly influence the specific aspect of disease addressed in the critical question.



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### Appendix E1. Literature classification schema.\*

Design/ Class	Therapy <sup>†</sup>	Diagnosis <sup>‡</sup>	Prognosis§
1	Randomized, controlled trial or meta-analysis of randomized trials	Prospective cohort using a criterion standard or meta-analysis of prospective studies	Population prospective cohort or meta-analysis of prospective studies
2	Nonrandomized trial	Retrospective observational	Retrospective cohort Case control
3	Case series	Case series	Case series

<sup>\*</sup>Some designs (eg, surveys) will not fit this schema and should be assessed individually.

**Appendix E2.** Approach to downgrading strength of evidence.

	Design/Class			
Downgrading	1	2	3	
None	ı	П	III	
1 level	II	III	X	
2 levels	III	X	X	
Fatally flawed	X	X	X	

Appendix E3. Likelihood ratios and number needed to treat.\*

LR (+)	LR (-)	
1.0	1.0	Does not change pretest probability
1–5	0.5–1	Minimally changes pretest probability
10	0.1	May be diagnostic if the result is concordant with pretest probability
20	0.05	Usually diagnostic
100	0.01	Almost always diagnostic even in the setting of low or high pretest probability

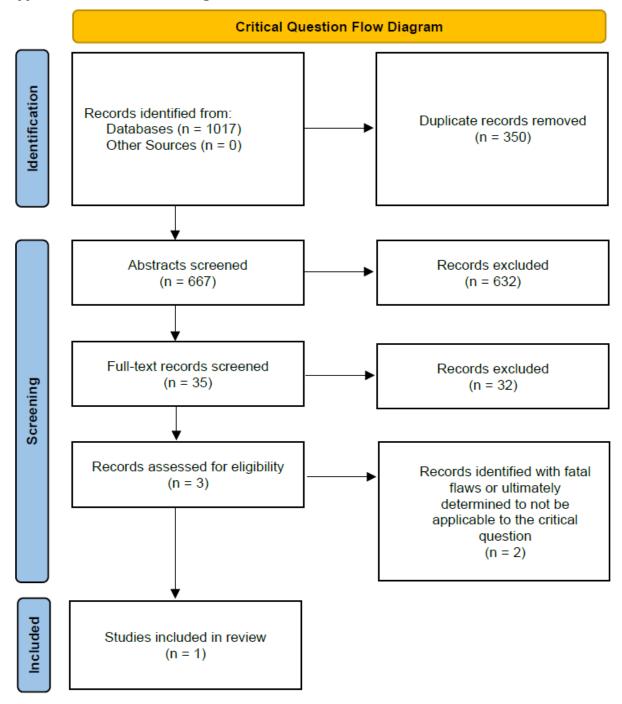
LR, likelihood ratio.

<sup>†</sup>Objective is to measure therapeutic efficacy comparing interventions.

<sup>&</sup>lt;sup>‡</sup>Objective is to determine the sensitivity and specificity of diagnostic tests.

<sup>§</sup>Objective is to predict outcome, including mortality and morbidity.

<sup>\*</sup>Number needed to treat (NNT): number of patients who need to be treated to achieve 1 additional good outcome; NNT=1/absolute risk reduction×100, where absolute risk reduction is the risk difference between 2 event rates (ie, experimental and control groups).



# **Appendix E5. Literature Searches**

Search Date	Database	Search Strings	Filters
8/23/2022	PubMed	(("Hypertension" [tiab]) OR ("Blood Pressure" [tiab]) OR ("Hypertension" [MH]) OR ("Blood Pressure" [MH])) AND (("Antihypertensive" [tiab]) OR ("Clevidipine" [tiab]) OR ("Enalaprilat" [tiab]) OR ("Esmolol" [tiab]) OR ("Fenoldopam" [tiab]) OR ("Hydralazine" [tiab]) OR ("Labetalol" [tiab]) OR ("Nicardipine" [tiab]) OR ("Nitroglycerin" [tiab]) OR ("Nitroprusside" [tiab]) OR ("Phentolamine" [tiab]) OR ("Antihypertensive Agents" [Pharmacological Action]) OR ("Clevidipine" [Supplementary Concept]) OR ("Enalaprilat" [MH]) OR ("Esmolol" [Supplementary Concept]) OR ("Fenoldopam" [MH]) OR ("Hydralazine" [MH]) OR ("Labetalol" [MH]) OR ("Nitroglycerin" [MH]) OR ("Nitroprusside" [MH]) OR ("Phentolamine" [MH]) AND (("Emergency Medicine" [tiab]) OR ("Emergency Treatment" [tiab]) OR ("Emergency Department" [tiab]) OR ("Emergency Medical Service*" [tiab]) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [tiab]) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH])) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Medical Services" [MH])) OR ("Emergency Treatment" [MH]) OR ("Emergency Treatment" [MH]) OR ("Em	2011- Current
8/24/2022	Scopus	TITLE-ABS-KEY("Hypertension" OR "Blood Pressure" OR "Hypertension") AND TITLE-ABS-KEY("Antihypertensive" OR "Antihypertensive Agent*" OR "Clevidipine" OR "Enalaprilat" OR "Esmolol" OR "Fenoldopam" OR "Hydralazine" OR "Labetalol" OR "Nicardipine" OR "Nitroglycerin" OR "Nitroprusside" OR "Phentolamine") AND TITLE-ABS-KEY("Emergency Medicine" OR "Emergency Treatment" OR "Emergency Department" OR "Emergency Medical Service*") AND NOT ("Pregnant" OR "Pregnancy" OR "Pregnancies") AND NOT ("Stroke") AND NOT ("Myocardial Ischemia") AND NOT ("Pulmonary Edemia") AND NOT ("Heart Failure")	2011- Current
8/24/2022	Embase	('asymptomatic':ti,ab,kw AND 'hypertension':de,ti,ab,kw OR 'pulmonary hypertension':de,ti,ab,kw) AND ('antihypertensive agent':de,ti,ab,kw OR 'antihypertensive therapy':de,ti,ab,kw OR 'clevidipine':de,ti,ab,kw OR 'enalaprilat':de,ti,ab,kw OR 'esmolol':de,ti,ab,kw OR 'fenoldopam':de,ti,ab,kw OR 'hydralazine':de,ti,ab,kw OR 'labetalol':de,ti,ab,kw OR 'nitroglycerin':ti,ab,kw OR 'glyceryl trinitrate':de,ti,ab,kw OR 'nitroprusside':ti,ab,kw OR 'nitroprusside'sodium':de,ti,ab,kw OR 'emergency medicine':de,ti,ab,kw OR 'emergency treatment':de,ti,ab,kw OR 'emergency department':ti,ab,kw OR 'emergency ward':de,ti,ab,kw OR 'emergency medical service*':ti,ab,kw OR 'emergency health service':de,ti,ab,kw) NOT ('Pregnant':ti,ab,kw OR 'Pregnancy':de,ti,ab,kw OR 'Pregnancies':ti,ab,kw) NOT ('Stroke':ti,ab,kw) NOT ('Mycoardial Ischemia':ti,ab,kw OR 'Heart Muscle Ischmeia':de,ti,ab,kw) NOT ('Pulmonary Edema':ti,ab,kw OR 'Lung Edema':de,ti,ab,kw) NOT ('Heart Failure':de,ti,ab,kw)	2011- Current

**Appendix E5. Literature Searches (continued)** 

Search Date	Database	Search Strings			
8/24/2022	Web of Science	TS=("Hypertension" OR "Blood Pressure" OR "Hypertension") AND TS=("Antihypertensive" OR "Antihypertensive Agent*" OR "Clevidipine" OR "Enalaprilat" OR "Esmolol" OR "Fenoldopam" OR "Hydralazine" OR "Labetalol" OR "Nicardipine" OR "Nitroglycerin" OR "Nitroprusside" OR "Phentolamine") AND TS=("Emergency Medicine" OR "Emergency Treatment" OR "Emergency Department" OR "EMS" OR "Emergency Medical Service*") NOT TS=("Pregnant" OR "Pregnancy" OR "Pregnancy" OR "Stroke" OR "Myocardial Ischemia" OR "Pulmonary Edemia" OR "Heart Failure")	2011- Current		
8/24/2022	Cochrane Library	('asymptomatic':ti,ab,kw AND 'hypertension':ti,ab,kw OR 'pulmonary hypertension':ti,ab,kw) AND ('antihypertensive agent':ti,ab,kw OR 'antihypertensive therapy':ti,ab,kw OR 'clevidipine':ti,ab,kw OR 'enalaprilat':ti,ab,kw OR 'esmolol':ti,ab,kw OR 'fenoldopam':ti,ab,kw OR 'hydralazine':ti,ab,kw OR 'labetalol':ti,ab,kw OR 'nitroglycerin':ti,ab,kw OR 'glyceryl trinitrate':ti,ab,kw OR 'nitroprusside':ti,ab,kw OR 'nitroprusside sodium':ti,ab,kw) AND ('discharge':ti,ab,kw OR 'patient discharge':ti,ab,kw OR 'hospital discharge':ti,ab,kw) AND ('emergency medicine':ti,ab,kw OR 'emergency treatment':ti,ab,kw OR 'emergency department':ti,ab,kw OR 'emergency ward':ti,ab,kw OR 'emergency medical service*':ti,ab,kw OR 'emergency health service':ti,ab,kw)	2011- Current		

Appendix E6. Quality Payment Program (QPP) **Measure ID** QPP317 Preventive Care and Screening: Screening for High Blood Pressure and Follow-Up Documented. **Measure Description** Percentage of patients aged 18 years and older seen during the submitting period who were screened for elevated blood pressure AND a recommended follow-up plan is documented based on the current blood pressure reading as indicated. **Data of Interest** Patients who were screened for elevated blood perssure have a recommended followup plan documented, as indicted if the blood pressure is pre-hypetentsive or hypertensive All patients aged 18 years and older at the beginning of the measurement period with at least one eligible encounter during the measurement period **Denominator Exclusions:** Patient not eligible due to active diagnosis of hypertension. **Denominator Exceptions:** Patient refuses to participate (either blood pressure measurement or follow-up). Patient is in an urgent or emergent medical situation where time is of the essence and to delay treatment would jeopardize the patient's health status. This may include but is not limited to severely elevated blood pressure when immediate medical treatment is indicated. Documented reason for not screening or recommending a follow-up for high blood pressure. **Numerator Exclusions:** Not Applicable 

Evidentiary Table.

<b>Author &amp; Year</b>	Class of	Setting &	Methods & Outcome	Results	Limitations & Comments
Published	Evidence	Study Design	Measures		
Brody et al (2015)	Ш	Secondary analysis of data pooled from two randomized controlled trials; single, urban, academic medical center	Included emergency department patients with asymptomatic hypertension and subclinical hypertensive heart disease; patients with uncontrolled blood pressure (>140/90 mm Hg in one study and >160/90 mm Hg for the other study) and discharged from the emergency department; excluded potential hypertensive emergencies or cardiovascular or neurovascular events; outcomes: short-term blood pressure reduction; adverse events; multivariable regression to evaluate association with antihypertensive initiation from the emergency department and blood pressure reduction	N=217; baseline characteristics were similar between those who received an antihypertensive prescription and those who did not except for higher systolic blood pressure among those who received a prescription; systolic blood pressure reduction was independently associated with antihypertensive prescriptions from the emergency department ( <i>P</i> =.001); the antihypertensive prescription accounted for a reduction of 11 mm Hg (95% CI 4 to 17 mm Hg; <i>P</i> =.001); adverse events were comparable and low in both groups	Retrospective; small number of observations from 1 health system; predominantly black population (96%)