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Hypertensive Disorders of Pregnancy



Innovative Management Strategies

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ABSTRACT

Hypertensive disorders of pregnancy (HDP) complicate 13% to 15% of pregnancies in the United States. Historically marginalized communities are at increased risk, with preeclampsia and eclampsia being the leading cause of death in this population. Pregnant individuals with HDP require more frequent and intensive monitoring throughout the antepartum period outside of routine standard of care prenatal visits. Additionally, acute rises in blood pressure often occur 3 to 6 days postpartum and are challenging to identify and treat, as most postpartum individuals are usually scheduled for their first visit 6 weeks after delivery. Thus, a multifaceted approach is necessary to improve recognition and treatment of HDP throughout the peripartum course. There are limited studies investigating interventions for the management of HDP, especially within the United States, where maternal mortality is rising, and in higher-risk groups. We review the state of current management of HDP and innovative strategies such as blood pressure self-monitoring, telemedicine, and community health worker intervention. (JACC Adv 2024;3:100864) © 2024 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

EPIDEMIOLOGY OF HYPERTENSIVE DISORDERS OF PREGNANCY

Hypertensive disorders of pregnancy (HDP), defined as preeclampsia or eclampsia, gestational hypertension, and chronic hypertension, complicate 13% to 15% of all pregnancies in the United States (Table 1).¹ Physicians and health care providers must be vigilant for the diagnosis across specialties. Criteria for diagnosis of hypertension in pregnancy include 2 readings of a systolic blood pressure (BP) \geq 140 mm Hg and/or a diastolic BP \geq 90 mm Hg, taken over a period of 4 to 6 hours. Preeclampsia, defined as an acute rise in BP and proteinuria (\geq 300 min/24 h or \geq 1+ with dipstick) after the 20th week of pregnancy or postpartum, affects approximately 2 to 8% of pregnant women.² Preeclampsia with severe features includes one or more of the following: a BP \geq 160/110 mm Hg, proteinuria (2.0 g/24 h or \geq 2+ dipstick), new-onset serum creatinine >1.2 mg/dL, platelets <100,000/mm³,

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ABBREVIATIONS AND ACRONYMS

BP = blood pressure

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HDP = hypertensive disorders of pregnancy increased lactate dehydrogenase, elevated alanine aminotransferase or aspartate aminotransferase, or persistent headache or other cerebral or visual disturbance and persistent epigastric pain. Preeclampsia can progress to eclampsia, additionally defined by

new-onset seizures during or after labor.³ Gestational hypertension is diagnosed as hypertension diagnosed for the first time during pregnancy that returns to normal <12 weeks postpartum and without proteinuria or diagnostic features of preeclampsia.⁴ Chronic hypertension is persistence of elevated BP \geq 12 weeks postpartum or hypertension in pregnancy that was also present prior to 20 weeks' gestation.⁴ Together, HDP accounts for 7% of maternal deaths in the United States.⁵ The incidence of preeclampsia has increased over recent decades, surpassing the rate of diseases like Alzheimer's, obesity, diabetes, and chronic kidney disease.^{6,7} The incidence of HDP has exponentially increased from 528.9 per 10,000 deliveries in 1993 to 912.4 per 10,000 deliveries in 2014.⁸ Risk factors for preeclampsia include age, with a higher incidence among those ages <25 and >35 years, diabetes, obesity, and preexisting cardiovascular disease (CVD).7 Additionally, dietary intake of sodium has been found to increase risk for HDP, with those consuming >3.5 g/d having a 54% higher risk for chronic hypertension and a 20% higher risk for preeclampsia compared to those consuming <2.8 g/d.⁸

HDP is prevalent and undertreated in pregnant individuals from historically marginalized communities due to systemic barriers to health care. There is increased risk for preeclampsia among minority groups, with higher frequency of preeclampsia among non-Hispanic Black women compared to non-Hispanic White women (16.7% vs 13.4%, respectively).^{8,9} Maternal mortality is approximately 3-fold higher in Black individuals, with preeclampsia and eclampsia leading causes of death, accounting for

HIGHLIGHTS

- HDP complicate 13 to 15% of all pregnancies in the United States.
- Individuals with HDP require more frequent and intensive peripartum management than routine standard of care.
- Innovative management strategies are needed, but the best approach, timing, frequency, and intensity are uncertain.
- Blood pressure self-monitoring, telemedicine, and community health worker intervention may be novel approaches to HDP.

30% of pregnancy-related deaths.⁹ A single-center prospective study that screened >10,000 postpartum women detected a new diagnosis of postpartum hypertension in 8%, most frequently in Black individuals with higher body mass index.¹⁰ Short-term adverse outcomes resulting from preeclampsia include increased risk for cesarean delivery, placental abruption, prolonged maternal hospital stay, and increased mortality. Those who develop preeclampsia prior to 37 weeks have an 8fold increase in adverse pregnancy outcomes like preterm birth and placental abruption.⁷ Additionally, hypertension is the foremost indication for postpartum readmission.¹¹

Long-term adverse outcomes due to HDP are well documented. Pregnant individuals with HDP have approximately double the risk of ischemic heart disease in the first 12 years after pregnancy compared to those who are normotensive.¹² There may be a significant relationship between the timing of onset of

TABLE 1 Definitions of	Hypertensive Disorders of Pregnancy
Chronic HTN	SBP \geq 140 mm Hg or DBP \geq 90 mm Hg on at least 2 occasions 4 h apart at a gestational age of $<$ 20 wk
Gestational HTN	SBP ≥140 mm Hg or DBP ≥90 mm Hg on at least 2 occasions 4 h apart at a gestational age of >20 wk with no features of preeclampsia
Preeclampsia	Without severe features: SBP ≥140 mm Hg or DBP ≥90 mm Hg on at least 2 occasions 4 h apart at a gestational age of >20 wk with proteinuria (>300 mg on 24-h urine protein collection or 0.3 mg on urine point of care)
	 With severe features: SBP ≥140 mm Hg or DBP ≥90 mm Hg on at least 2 occasions 4 h apart at a gestational age of >20 wk with proteinuria and evidence of end organ injury SBP ≥160 mm Hg or DBP ≥110 mm Hg on at least 2 occasions 4 h apart OR persistent severe hypertension requiring IV antihypertensives for control to bring SBP <160 mm Hg or DBP <110 mm Hg
Eclampsia	Preeclampsia + seizures
DBP = diastolic blood pressure	; HTN = hypertension; SBP = systolic blood pressure.

preeclampsia and long-term cardiovascular outcomes. Individuals who develop preeclampsia after 37 weeks have a 2-fold increase in long-term adverse cardiovascular outcomes.7 There are data demonstrating an association of advanced maternal age (35-44 years) and very advanced maternal age (>45 years) on HDP risk itself, though whether that translates to increased risk of major cardiovascular events in older women is not well established.13 Individuals with HDP have a higher risk of developing stroke compared to those who do not have HDP (34.5% vs 6.9%), and the risk of stroke is higher among non-Hispanic Black and Hispanic/Latina women as compared to non-Hispanic White women.⁸ Preeclampsia is also associated with adverse longterm kidney outcomes, including an increased risk of developing glomerular or proteinuric kidney disease and end-stage kidney disease within 5 to 10 years after pregnancy.¹⁴ The risk is higher in women with multiple preeclamptic pregnancies or previous preterm preeclampsia.

Late-onset preeclampsia can present in individuals with or without history of antepartum HDP. Acute BP elevation is frequently observed 3 to 6 days postpartum. Thus, the standard of care postpartum visit at 6 weeks after delivery fails individuals with preeclampsia occurring in the first few days postpartum. Most readmissions for acute hypertension occur within 10 to 20 days of discharge, well before the usual postpartum visit.¹⁵ From the American College of Obstetricians and Gynecologists recommendations, "blood pressure evaluation is recommended for women with hypertensive disorders of pregnancy no later than 7 to 10 days postpartum, and women with severe hypertension should be seen within 72 hours; other experts have recommended follow-up at 3 to 5 days".¹⁶ Prioritizing those with the highest BPs and more severe features for postpartum BP checks or visits by 3 days appears reasonable.

Individuals without a previous diagnosis of hypertension are at particularly increased risk for severe maternal morbidity from acute BP elevation compared to those with chronic hypertension. Over half of all maternal deaths occur in the postpartum period.¹⁷ Causes of death from HDP include systolic heart failure, cerebrovascular disease, myocardial infarction, and cardiac arrest.¹⁸ In the context of HELLP syndrome (hemolysis, elevated liver enzymes, and low platelets), additional causes of HDP-related death are placental abruption, acute respiratory distress syndrome, disseminated intravascular coagulation, hepatic hemorrhage, hypoxic ischemic encephalopathy, and acute kidney injury.¹⁹ Among HDP-related deaths, 44.3% of deaths occurred on day

1 and 37.1% of deaths on days 2 to 7.²⁰ In a review of 232 pregnancy-related deaths evaluated by 13 state Maternal Mortality Review Committees, approximately 60% of postpartum maternal deaths were determined to be preventable.²¹ Prevention of mortality likely requires a multipronged approach from highly responsive health systems, such as patient education on BP, healthy diet/lifestyle, medical therapy (aspirin, antihypertensives, magnesium), surveillance for proteinuria and elevated BP, use of safety bundles, strategically timed delivery, and provider action to treat emergent symptoms.^{22,23} The immediate postpartum period, commonly termed the "fourth trimester," is a prime window of opportunity for intervention and transition to primary or cardiovascular care. Short-term management of HDP immediately postpartum may influence long-term BP control, although data is limited in this regard.²⁴

PATHOPHYSIOLOGY OF HYPERTENSIVE DISORDERS OF PREGNANCY

The pathophysiology of HDP is not well understood. Hypertension in pregnancy is thought to be due to a combination of improper trophoblast differentiation and abnormal regulation of cytokines, adhesion molecules, major histocompatibility complex molecules, and metalloproteinases, which lead to abnormal differentiation of spiral arteries and subsequent placental hypoperfusion and ischemia.²⁵ Antiangiogenic factors that are released also play a role in the development of systemic hypertension due to systemic endothelial dysfunction.²⁵ Current research includes several hypotheses as to how preeclampsia can occur, including contact between the maternal immune system and the placental semiallogeneic trophoblast, chronic uteroplacental ischemia, oxidative stress, immune maladaptation, imbalance of angiogenic factors, genetic imprinting, altered renal hemodynamics with impaired renin angiotensin signaling, an exaggerated maternal inflammatory response, and vascular endothelial dysfunction.²⁶

Angiogenic factors play a key role in regulation of placental vascular differentiation; thus, there may be an imbalance between angiogenic factors in the pathogenesis of preeclampsia.²⁷ In early gestation of normal pregnancy, proangiogenic factors like endoglin (Eng), placental growth factor (PIGF), fms-like tyrosine kinase-1(Flt1), and vascular endothelial growth factor are highly expressed by invasive trophoblasts. However, decreased expression of these factors can lead to the inadequate cytotrophoblastic invasion seen in preeclampsia. The splice variants of

Flt1 and Eng, namely soluble Flt1 (sFlt1) and soluble Eng (sEng), serve as ligand traps by binding to angiogenic factors. In normal pregnancy, sFlt-1 blood levels remain low in early pregnancy and increase toward the third trimester to allow for cytotrophoblastic invasion during early pregnancy. However, elevated levels of sFlt1 in pregnancy, as seen in preeclampsia, can lead to defective cytotrophoblast invasion and high plasma sFlt1:PlGF ratios, reflecting severe disease and associated with adverse clinical outcomes. Levels of sFLT-1 can persist early postpartum and lead to preeclampsia usually within 48 hours to 6 weeks postpartum. Interestingly, postpartum preeclampsia has also been associated with epigenetic modifications, with CpG methylation sites as markers for identification of postpartum preeclampsia.27,28

CURRENT MANAGEMENT OF HYPERTENSIVE DISORDERS OF PREGNANCY

Preventing HDP altogether is most optimal to avoid adverse outcomes. There appears to be an inverse relationship between physical activity, gestational hypertension, and preeclampsia.²⁹ Preventative medical management with aspirin is recommended per American College of Obstetrics and Gynecology (ACOG) guidelines for those at increased risk for preeclampsia, including those with previous preeclampsia, multifetal gestation, chronic kidney disease, autoimmune disease, type 1 or type 2 diabetes mellitus, and chronic hypertension.³⁰ For these individuals, there is a Level of Evidence: A recommendation to start low-dose aspirin (81 mg) before 16 weeks of gestation for prevention of preeclampsia. Individuals who develop preeclampsia after 37 weeks of gestation are recommended to undergo immediate delivery.^{22,30}

Severe hypertension and preeclampsia must be treated early on to prevent adverse outcomes. The BP target for acute severe hypertension and preeclampsia ante or postpartum is a BP of 140-150/90-100 mm Hg.³¹ The optimal BP target for the treatment of mild chronic hypertension is an area of active investigation. In an open-label, multicenter, randomized controlled trial (RCT) assigning pregnant women with mild chronic hypertension (BP <160/100 mm Hg) to antihypertensive medication, a BP target of 140/90 mm Hg in patients with chronic hypertension was associated with lower risk of developing preeclampsia with severe features, medically indicated preterm birth (<35 weeks of gestation), placental abruption, or fetal or neonatal death, and there was no increase in the risk for small-for-gestational-age birth weight.³² Medical therapies for HDP antepartum include hydralazine, labetalol, nifedipine, and methyldopa. Postpartum, the angiotensin-converting enzyme inhibitor, enalapril, may be useful and is compatible with breastfeeding in normal-term neonates.³³ Magnesium is used for seizure prophylaxis and diuretics for edema or volume overload postpartum.

The routine use of diuretics is unclear in preeclampsia.³⁴ Valensise et al³⁵ assessed pregnant individuals with uterine artery Doppler to evaluate placental arterial waveforms and maternal transthoracic echocardiography, calculating stroke volume and peripheral vascular resistances. At a hemodynamic level, there appeared to be 2 phenotypes of preeclampsia: 1) early onset (<34 weeks), characterized by low cardiac output, high resistance, and depleted intravascular volume, a phenotype more commonly associated with bilateral notching of the uterine artery Doppler, fetal growth restriction, and worse maternal and perinatal outcomes; and 2) late onset (\geq 34 weeks), characterized by high cardiac output, reduced resistance, and increased intravascular volume, a phenotype more commonly associated with obesity, normal fetal growth, and more favorable maternal and perinatal outcomes. Understanding different preeclampsia phenotypes could modify the choice of therapy, specifically the use of diuretics.

Following delivery, fluid that has been sequestered in the extravascular space is mobilized, producing a large auto-infusion of fluid from the extravascular to the intravascular compartment. Trials in antepartum patients have shown insufficient evidence to draw reliable conclusions about diuresis, possibly because these could have also included the phenotype associated with intravascular depletion.^{35,36} Several studies have demonstrated that diuretics may be useful in postpartum women with HDP, possibly including more individuals with a phenotype of increased intravascular volume. Small trials of individuals with severe preeclamptic features postpartum have shown decreased requirement for additional antihypertensives if a combination of furosemide and nifedipine was used.³⁷ Patients with preeclampsia with severe features randomized to treatment with 20 mg daily furosemide were found to have significantly lower BP by postpartum day 2 and required significantly less antihypertensive therapy on discharge compared to those treated with placebo.³⁸ Another study evaluating patients with gestational hypertension and preeclampsia with and without severe features demonstrated that patients randomized to furosemide 20 mg daily for the first 5 days postpartum were less likely to have persistent

hypertension at postpartum day 7.³⁹ Additional research is needed to validate different preeclampsia phenotypes and the best therapy.

Given that preeclampsia can develop de novo after delivery, women in the postpartum period should be given discharge instructions that include information about the signs and symptoms of preeclampsia as well as the importance of presenting for medical evaluation in the event that they occur.¹⁶ As delineated by the ACOG guidelines, treatment with first-line agents in patients with acute severe hypertension, identified as systolic BP ≥160 mm Hg or diastolic BP \geq 110 mm Hg, should be initiated within 30 to 60 minutes to reduce risk of maternal stroke. First-line agents for treatment in both pregnancy and postpartum include IV labetalol and hydralazine, or more recently, the use of oral nifedipine if IV access is not established. BP targets in cases of severe hypertension should not aim for normalization of BP but rather target a range of 140 to 150/90 to 100 mm Hg. In severe cases, such as when medication fails to reduce BP to the target range, consultation with anesthesiologist, maternal fetal medicine subspecialist, or critical care subspecialist is recommended.³¹

ACOG and the Royal College of Obstetricians and Gynecologists recommend achieving a BP of 140/90 mm Hg in the immediate postpartum period.^{40,41} However, there are no standardized management guidelines for specific antihypertensive agents or parameters for medication uptitration in the postpartum period. Thus, physician preference and experience, as well as safety of medical therapy during breastfeeding often affect the approach.⁴²

INNOVATIVE MANAGEMENT OF HYPERTENSIVE DISORDERS OF PREGNANCY

BLOOD PRESSURE SELF-MONITORING AS A STRATEGY FOR TREATMENT OF HYPERTENSIVE DISORDERS OF PREGNANCY. Self-monitoring with a health systemprovided BP cuff, when combined with outreach to patients by medical personnel for antihypertensive management, appears to be a viable strategy for antepartum BP control (Table 2). Two well-powered RCTs, BUMP1 and BUMP2, did not find differences between those randomized to self-monitoring of BP vs usual care.43,44 However, those trials did not include automated transfer of BP readings to physicians via the electronic medical record (EMR) or outreach by the medical team; instead, they relied on the volunteer to contact the medical team for elevated BP readings. An observational study studying universal screening noted lack of participation in

self-monitoring unless a BP cuff was provided.¹⁰ By comparison, SNAP-HT demonstrated feasibility and improved diastolic BP in the intervention group at 6 months with a 7 mm Hg decrease in long-term BP using self-management of postpartum hypertension with daily home BP monitoring and automated selfcontrolled medication adjustment via telehealth.45 High volunteer satisfaction was noted in 4 studies: a qualitative analysis of the BUMP2 study, Heart Safe Motherhood, a University of Wisconsin study, and Safe@Home.44,46-48 In these studies, there were frequent reminders and contact with the study teams. In BUMP2, volunteers highlighted interactions with clinicians, structured follow-up, and individualized support as aspects that they preferred and were motivated to reduce HDP in future pregnancies. There were significant health knowledge gaps despite college-level education, pointing to the importance of universal education on HDP.44 In the University of Wisconsin study of postpartum management, where telehealth was used for communication, there was a 95% retention rate and high patient satisfaction, and investigators also identified 16% of their cohort with uncontrolled BP.47 The Safe@Home study using telemonitoring to monitor BP, admissions for suspected preeclampsia and hypertension were lower in those with telemonitoring compared to those without, in addition to high levels of satisfaction in the telemonitoring group.⁴⁸ Additionally, an observational study investigating self vs ambulatory BP monitoring, found that patients were more comfortable with selfmonitoring, citing less anxiety and discomfort.49 Furthermore, a University of Pittsburgh observational study found that when patients were given educational materials on discharge, with follow-up at 1-week intervals, studying universal screening noted lack of participation in self-monitoring; in those who did participate, 8% of the cohort was diagnosed with potential new-onset hypertension and 0.7% were diagnosed with severe hypertension.¹⁰ Interestingly, a meta-analysis of home monitoring showed no significant differences between those as compared to usual care with respect to postpartum readmission and BP monitoring acutely after discharge, possibly due to clinical heterogeneity and low quality of evidence.⁵⁰ While more RCT data are needed, a multilevel intervention with patient-provider interaction appears more effective in achieving the goal of BP reduction than self-monitoring in isolation.

TELEMEDICINE FOR HYPERTENSIVE DISORDERS OF PREGNANCY. Greater than 40% of women do not attend a recommended postpartum visit by 6 weeks; thus, it is important to study the utilization of

TABLE 2 Rigor and	Reproducibility of Random	nized Controlled Trials and	d Observational Studies of Blood P	ressure Monitoring in Per	ipartum Cohorts
First Author, Study	Sample	Design	Intervention	Follow-up Time	Key Findings
Tucker et al, ⁴³ BUMP 1	Antepartum, England, >70% White	RCT N = 2,441	Self-monitoring, telemonitoring, prompt to contact clinic for high BP	Until Delivery	No change in time to first clinic recording of HTN by provider
Chappell et al, ⁴⁴ BUMP2	Antepartum, England, >49% White	$\begin{array}{l} RCT \\ N = 850 \end{array}$	Self-monitoring, telemonitoring, prompt to contact clinic for high BP	Until Delivery	No change in mean SBP
Cairns et al, ⁴⁵ SNAP-HT	Postpartum, England, >80% White	RCT N = 91	Self-monitoring, automated medication adjustment via telemonitoring + home visits	6 mo	90% completion SBP and DBP lower in intervention group most markedly at 6 wk, DBP lower at 6 mo
Janssen et al, ⁴⁶ Heart Safe Motherhood	Postpartum, U.S. ~42% White, 34% Black, 12% Hispanic/Latina	Observational N = 199	Text based informing of BP, physician-directed therapy or admission	Postpartum day 10	97% of participants submitted 1 BP
Hoppe KK et al ⁴⁷	Postpartum, U.S. 93% White, 9.1% Hispanic/Latina	Prospective single cohort study N = 55	Telehealth intervention for BP monitoring	6 wk postpartum	Retention rate 95%, incidence of severe HTN was 16%, 53% of participants required treatment, no readmissions, 86% of participants satisfied
Van den Heuvel et al, ⁴⁸ Safe@Home	Antepartum, the Netherlands, >75% White	Case-control N = 236	Self-monitoring, provider acting on automated results	16 wk gestational age continued until delivery	Less antenatal visits and admission in intervention group, high satisfaction
Hacker et al ¹⁰	Postpartum, U.S. >67% non- Hispanic White	Prospective observational N = 1,192	Given educational materials on discharge, follow-up with nurse at 1-wk intervals	NA	26% could not participate because they did not have BP cuff. Of the remainder participating, 8% had a potential new diagnosis of a hypertensive disorder of pregnancy, and 0.7% having severe hypertension
Taylor et al ⁴⁹	Antepartum and postpartum, Auckland New Zealand	Observational N = 113	Given self BP cuff or ambulatory cuff (24-h monitor)	24 h	Most participants preferred self BP checks, citing ambulatory cuff caused more anxiety, discomfort
DBP = diastolic blood pres	ssure; ED = emergency departme	ent; HTN = hypertension; NA =	not available; RCT = randomized controlled	trial; SBP = systolic blood pres	sure; SPEC = severe preeclampsia.

telehealth and telemedicine to aid in attendance of these visits.¹⁶ Telemedicine is a viable alternative with greater attendance to postpartum visits in several studies (Table 3). A single center study in a U.S. urban minority community indicated women who did not attend an in-person office visit were more often Black (87% vs 56%), P < 0.01) and younger (29.1 vs 31.4 years, P = 0.04), but no disparity by race or ethnicity was seen for telehealth visits.⁵¹ Attendance for a telehealth visit was high, 70% vs 32% for an in-person visit.⁵¹ Telephone, text message, and video conference-based communication were well received by patients and providers in several studies.^{52,53} However, a University of Arkansas study of self-monitoring in a rural cohort using mobile health (mHealth) wireless transmitting equipment found several challenges with this strategy.⁵² Those using the application found it easy to use, while nonusers were concerned about incorporating it into their daily routine as new parents. Barriers to using mHealth included concern for wireless transmission in rural areas, single BP cuff size availability, and stress associated with monitoring of BP. Most telehealth or mobile health studies with BP cuff prototypes have been tested in small studies and lack integration with EMRs or broader medical practice.⁵⁴ The "OB Nest" program at Mayo Clinic is an antepartum model involving self-monitoring, texting, and an online peer support community.⁵³ The program has reduced in-person prenatal care, decreasing health care costs, with similar delivery outcomes. Participants in the program reported feeling more

TABLE 3 Rigor and	l Reproducibility of Pre	vious Studies of Teleme	dicine in Peripartum Coh	iorts	
First Author, Study	Sample	Design	Intervention	Follow-Up Time	Key Findings
Sanghavi et al ⁵¹	Postpartum, U.S., >73% Black	Retrospective cohort N = 119	Telemedicine and in-person	NA	70% completion for telemedicine vs 32% for in-person, those who were more likely to not complete an in-person visit were more likely to be Black (87% vs 56%) and younger (29.1 vs 31.4); this difference not seen with telemedicine
Payakachat et al, ⁵² mHealth	Postpartum, U.S., >48% White	Prospective cohort N = 37	Self-monitoring of BP, weight, pulse, O ₂ sat, and symptom survey with wireless transmission to providers	NA	Users had higher facilitating condition scores, higher levels of perceived benefits, and lower levels of perceived barriers vs nonusers, increased admission in users
De Mooji et al, ⁵³ OB Nest	Antepartum (<5 mo gestation), U.S., 85% White	Observational study $N = 20$	Self-monitoring tools, online communities, and text-based communication	Antepartum to 6 wk postpartum	Improved patient satisfaction, lower anxiety levels, lower health care costs, no difference in delivery outcomes
Hirshberg et al ^{55,56}	Postpartum, U.S., >66% Black	RCT N = 206	Self-monitoring of BP with text messaging platform	3 wk after delivery	Increase in at least 1 BP measurement in the first 10 d postpartum in the texting; greater attendance (>90%) to postpartum visits vs usual care among Black patients
Khosla et al ⁵⁷	Postpartum, U.S., >74% non- Hispanic Black	Retrospective cohort $N = 473$	Telehealth visits during the COVID pandemic to replace in person standard postpartum visit.	6 wk after delivery	Increased attendance of follow-up appointment in telehealth group vs in person, especially in Black individuals; Race-ethnic gap in care eliminated using telehealth
Rossiter et al ⁵⁸ , BP2	Postpartum, Australia, >64% White	RCT N = 157, N = 34 interviewed	Three arms: 1) usual care; 2) brief education intervention; 3) extended lifestyle intervention, 1 and 2 + phone based service, dieticians, exercise physiologists	12 mo postpregnancy	Extended lifestyle intervention increased recognition of their cardiac health risk, with greater motivation to make lifestyle and behavioral changes
Rich-Edwards et al, ⁵⁹ Heart Health 4 Moms (HH4H)	Postpartum, U.S., mostly White	RCT N = 151	Online educational modules, community forum and peer coaching vs control (internet links alone)	9 mo	Intervention participants reported significantly greater knowledge of cardiovascular risk factors, healthy eating, more physical activity vs controls
Herring et al ⁶⁰	Postpartum, U.S., 100% Black or Hispanic/Latina	RCT N = 22	Peer coach by phone, in-person, social media interaction	14 wk after delivery	Increased weight loss among intervention vs usual care
NA = Not applicable; BP	= blood pressure; RCT = ran	domized controlled trial.			

connected with providers, less anxious, and more knowledgeable.⁵³ A randomized control trial at the University of Pennsylvania used texting to communicate postpartum BP.⁵⁵ Their study found that the texting group was more likely to identify BP spikes than those presenting to office visits alone in the first 10 days postpartum, a high-risk period.⁵⁵ Use of telehealth may also reduce racial disparities.^{51,56,57} In a study at the University of Pennsylvania, Hirshberg et al⁵⁶ found decreased racial disparity when using a text-messaging-based platform. Their study found that over 90% of Black participants provided a BP measurement compared to the 33% who presented for an in-person BP visit as

per routine care. The implementation of telehealth with audio-based visits in a study done by Khosla et al⁵⁷ found significant improvement in adherence with at least 1 visit for follow-up for hypertension in the postpartum period among Black patients (48.5%-76.3%, P < 0.001).

Lifestyle interventions have been conducted via televisits. In a subset of the BP2 trial, women were more likely to adopt healthy habits if provided with an intense intervention with in-person consultation with a provider and dietitian followed by 6 months of telephone-based coaching.⁵⁸ Participants cited high accountability, increased education about healthy habits, and more motivation toward making lifestyle changes along with increased perceived risks of their cardiovascular health postpregnancy.⁵⁸ In the HH4H (Heart Health 4 Moms) study, a total of 150 women were enrolled in an RCT to reduce CVD risk through an online intervention vs self-directed care in controls.⁵⁹ The HH4H group improved CVD risk knowledge, self-efficacy to achieve a healthy diet, and reduced physical inactivity. This study enrolled mostly White, higher-income, and college-educated women who were normotensive at baseline.59 A small RCT in urban minority centers (Philadelphia Women, Infants, and Children program) found the use of texting BP measurements to providers and peer coaching to improve the frequency of BP measurement and weight loss.⁶⁰ Though data from rigorous RCTs are limited, telehealth may improve patientprovider communication, overcome caregiving and logistical barriers to in-person visits, and extend medical care into the home.

COMMUNITY HEALTH WORKERS FOR PATIENT SUPPORT AND EDUCATION IN HYPERTENSIVE DISORDERS OF **PREGNANCY.** The philosophy behind community health work is to recruit and train individuals with shared lived experiences to provide supportive care to patients either in the community or within health systems. Therefore, community health workers (CHWs) often have congruent cultural and language identities with the community they serve. CHWs provide peer support and linkage to health care and social services, extending the care team into homes across a region. They often provide practical information about exercise, diet, smoking cessation, and stress reduction, present information in layperson's terms, and their work may help in building a trusting relationship between the health system and community. Training is variable, with some programs offering core curricula and certifications. CHWs have most often been deployed in low- and middle-income countries or other resource-limited environments, but the Centers for Disease Control and American Public Health Association endorse the use of CHWs in chronic disease management, antiretroviral therapy for HIV, cancer screening, and asthma control.⁶¹

Few studies have focused on CHWs role in maternal health. The largest, the CLIP RCTs, deployed CHWs throughout India, Pakistan, and Mozambique for antepartum treatment of preeclampsia. The study was principally focused on the association of BP control with neonatal outcomes.⁶² CHWs were deployed to homes and used a digital phone-based application for patient assessment, measured BP, administered a dipstick urinalysis, gave patients oral and intramuscular medications, and referred patients for further care. The study found that CHWs were adequately utilized and that worsened BP control was associated with worse neonatal outcomes than for women with a normal BP but did not focus specifically on maternal health outcomes.63 The antepartum Health Start cohort in Arizona was a retrospective, propensity score-matched study of antepartum care in which patients were assigned a CHW. Involvement of a CHW increased prenatal care attendance significantly in a historically marginalized community of Latina and Native American individuals and mostly publicly insured cohort.⁶⁴ Though not a pregnancy-related study, CHW and telehealth were combined in the DREAM-GLOBAL study.65 A Canadian Indigenous and Tanzanian sample of middle-aged men and women involved a CHW to help volunteers measure BP on a centrally located device in a community health center, where they submitted measurements to a central server. Participants felt better educated on BP and lifestyle changes and perceived improved communication with physicians and support from site visits. CHW and health care providers found improved communication with participants and between each other.

CHWs have improved outcomes for individuals living with cardiovascular conditions (heart disease, stroke, type II diabetes), asthma, and cancer. A systemic review (including 16 RCTs but excluding maternal health) indicated a statistically significant reduction in emergency department (23%-51%), urgent care visits (60%), and hospitalization (21%-50%) when CHWs were integrated into the inpatient care team.⁶⁶ Combining CHWs with other strategies may be effective to reduce health care utilization but requires further investigation, particularly for antepartum or postpartum care. There are limited RCTs

Pre-pregnancy	 Identify patients at risk (age>35 years, prior preeclampsia, chronic hypertension, diabetes, thrombophilia, autoimmune disease, obesity, smoking) Optimize maternal health before pregnancy Increase awareness and empowering women regarding their health Teach self-care strategies
Antepartum	 Establish medical care with regular ObGyn visits and utilization of telehealth Self, in-person and remote monitoring of BP Education by and intervention from community health workers Identify and risk stratify patients who will benefit from additional monitoring Engage with peers and support groups
	1. Education and patient-provider dialogue regarding regular follow-up
Postpartum	 Follow-up (BP, medication and symptom review) Continuing self, in-person and/or remote monitoring Postpartum visits Transition to primary care Specialist follow-up

focused on cost and utilization with variability in the magnitude of the effect, likely due to differences in study design, and evidence is insufficient to draw conclusions about the effects of CHWs on chronic disease management.⁶⁷ A systematic review encompassing 8 studies (n = 6,500) found that 7 studies found no impact on mental health quality of life or mental health outcomes; 2 studies in the United States found improved quality of care in those with multiple-morbidity conditions and reduced hospitalizations.⁶⁸ However, these studies have low certainty of evidence due to risk of bias, inconsistency, and imprecision, thus pointing to a need for future studies investigating the role of CHWs in these conditions.⁶⁸ Important considerations for future studies of CHW include: tailored scope of work, training, mentorship, supervision, ratio of CHW to patient, and financing. The National Academies of Sciences, Engineering, and Medicine 2019 report provides health systems with guidance on how social care integration may promote improved health outcomes.⁶⁹

INNOVATING MULTILEVEL AND MULTIDISCIPLINARY MANAGEMENT OF HYPERTENSIVE DISORDERS OF PREGNANCY. A combination of strategies prepregnancy, antepartum, and postpartum can identify high-risk groups and patients at risk for developing HDP (Figure 1). Phone-based applications, blue-tooth or cellular upload of BP measurements to the EMR from BP self-monitoring, video-based EMR-integrated telemedicine, and CHWs could be integrated more fully to manage peripartum HDP (Central Illustration). Technology advances personcentered clinical care to increase health care access and empowers patients to engage in selfmanagement for improved health outcomes. A model of prioritized health care access that meets each patient where they are as opposed to relying on the patient coming to the health system may be more useful to identify acutely uncontrolled BP both ante and postpartum. CHWs can provide enhanced health education and coaching, peer support, and streamline patient-provider communication, aiding therapeutic optimization. CHWs could be incorporated into the cardio-obstetrics team, as consensus guidelines encourage early involvement of the cardio-obstetrics team to prevent maternal morbidity and mortality.⁷⁰ Furthermore, an equity-focused approach is crucial, with dedicated CHWs responsible for overseeing and encouraging retention in routine care, completion of referrals to primary and specialty care postpartum, screening for social needs, and introducing community resources. Assessment of clinical and social needs and coordination of real-time care plan management will allow for earlier identification, timely intervention, and prevention of unnecessary readmission and use of emergent/hospital-based care, as well as morbidity from uncontrolled BP.

Combined interdisciplinary approach for patients with HDP has been found effective.⁷¹ Interventions might include education for both providers discharging patients with HDP and patients with a diagnosis of HDP (by nurse educators or CHWs), the provision of free BP monitors to all patients with HDP,



scheduling of postpartum appointments prior to discharge, creation of a dedicated postpartum hypertension clinic, development of workflows and algorithms for those with HDP who present to the emergency department for evaluation, and potential readmission and development of workflows and algorithms for management of medication during the postpartum period. Implementation of this workflow and interventions can be associated with an increase in adherence to postpartum visit for follow-up for HDP.⁷¹

CONCLUSIONS

Combining elements of innovative management strategies like automated BP self-monitoring,

telemedicine, and individualized care by CHW with an interdisciplinary and equity-focused approach could be the future in managing HDP. If successful, this model could be generalized across centers and across other aspects of care, with the overall goal of reducing maternal morbidity and mortality.

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REFERENCES

1. Ford ND, Cox S, Ko JY, et al. Hypertensive disorders in pregnancy and mortality at delivery hospitalization - United States, 2017-2019. *MMWR Morb Mortal Wkly Rep.* 2022;71:585-591.

2. Johnson JD, Louis JM. Does race or ethnicity play a role in the origin, pathophysiology, and outcomes of preeclampsia? an expert review of the literature. *Am J Obstet Gynecol.* 2022;226: S876-s885.

3. Upadya M, Rao ST. Hypertensive disorders in pregnancy. *Indian J Anaesth*. 2018;62:675-681.

4. Wilkerson RG, Ogunbodede AC. Hypertensive disorders of pregnancy. *Emerg Med Clin.* 2019;37: 301–316.

5. Sharma G, Hays AG, Blumenthal RS. Can we reduce premature mortality associated with hypertensive disorders of pregnancy?: a window of opportunity. *J Am Coll Cardiol.* 2021;77:1313–1316.

6. Opichka MA, Rappelt MW, Gutterman DD, Grobe JL, McIntosh JJ. Vascular dysfunction in preeclampsia. *Cells.* 2021;10:3055.

7. Shih T, Peneva D, Xu X, et al. The rising burden of preeclampsia in the United States impacts both maternal and child health. *Am J Perinatol.* 2016;33:329-338.

8. Tsao CW, Aday AW, Almarzooq ZI, et al. Heart disease and stroke statistics-2022 update: a report from the American Heart Association. *Circulation*. 2022;145:e153-e639.

9. Cortés YI, Breathett K. Addressing inequities in cardiovascular disease and maternal health in Black women. *Circ Cardiovasc Qual Outcomes*. 2021;14:e007742.

10. Hacker FM, Jeyabalan A, Quinn B, Hauspurg A. Implementation of a universal postpartum blood pressure monitoring program: feasibility and outcomes. *Am J Obstet Gynecol MFM*. 2022;4:100613.

11. Clapp MA, Little SE, Zheng J, Kaimal AJ, Robinson JN. Hospital-level variation in post-partum readmissions. *JAMA*. 2017;317:2128-2129.

12. Bellamy L, Casas JP, Hingorani AD, Williams DJ. Pre-eclampsia and risk of cardiovascular disease and cancer in later life: systematic review and meta-analysis. *BMJ*. 2007;335:974.

13. Smithson SD, Greene NH, Esakoff TF. Pregnancy outcomes in very advanced maternal age women. *Am J Obstet Gynecol MFM*. 2022;4: 100491.

14. Vikse BE, Irgens LM, Leivestad T, Skjaerven R, Iversen BM. Preeclampsia and the risk of endstage renal disease. *N Engl J Med.* 2008;359: 800–809.

15. Wen T, Wright JD, Goffman D, et al. Hypertensive postpartum admissions among women without a history of hypertension or preeclampsia. *Obstet Gynecol.* 2019;133:712-719.

16. ACOG Committee opinion no. 736: optimizing postpartum care. *Obstet Gynecol.* 2018;131:e140-e150.

17. Declercq E, Zephyrin L. Maternal Mortality in the United States: A Primer. New York, NY: The Commonwealth Fund; 2020.

18. Ackerman CM, Platner MH, Spatz ES, et al. Severe cardiovascular morbidity in women with hypertensive diseases during delivery hospitalization. *Am J Obstet Gynecol*. 2019;220:582.e1-582. e11.

19. Isler CM, Rinehart BK, Terrone DA, Martin RW, Magann EF, Martin JN Jr. Maternal mortality associated with HELLP (hemolysis, elevated liver enzymes, and low platelets) syndrome. *Am J Obstet Gynecol.* 1999;181:924–928.

20. Dol J, Hughes B, Bonet M, et al. Timing of maternal mortality and severe morbidity during the postpartum period: a systematic review. *JBI Evid Synth*. 2022;20:2119-2194.

21. Petersen EE, Davis NL, Goodman D, et al. Vital signs: pregnancy-related deaths, United States, 2011-2015, and strategies for prevention, 13 states, 2013-2017. *MMWR Morb Mortal Wkly Rep.* 2019;68:423-429.

22. Davidson KW, Barry MJ, Mangione CM, et al. Aspirin use to prevent preeclampsia and related morbidity and mortality: US Preventive Services Task Force recommendation statement. *JAMA*. 2021;326:1186-1191.

23. Burgansky A, Montalto D, Siddiqui NA. The safe motherhood initiative: the development and implementation of standardized obstetric care bundles in New York. *Semin Perinatol.* 2016;40: 124–131.

24. Bramham K, Nelson-Piercy C, Brown MJ, Chappell LC. Postpartum management of hypertension. *BMJ*. 2013;346:f894.

25. Luger RK, Kight BP. *Hypertension in Pregnancy*. StatPearls; 2023.

26. Williams PJ, Broughton Pipkin F. The genetics of pre-eclampsia and other hypertensive disorders of pregnancy. *Best Pract Res Clin Obstet Gynaecol.* 2011;25:405-417.

27. Goel A, Maski MR, Bajracharya S, et al. Epidemiology and mechanisms of de novo and persistent hypertension in the postpartum period. *Circulation*. 2015;132:1726-1733.

28. Kim SK, Vishweswaraiah S, Macknis J, et al. New-onset postpartum preeclampsia: epigenetic mechanism and prediction. *J Matern Fetal Neonatal Med.* 2022;35:7179-7187. **29.** Dipietro L, Evenson KR, Bloodgood B, et al. Benefits of physical activity during pregnancy and postpartum: an Umbrella review. *Med Sci Sports Exerc.* 2019;51:1292–1302.

30. Gestational hypertension and preeclampsia: ACOG practice bulletin summary, number 222. *Obstet Gynecol.* 2020;135:1492-1495.

31. Committee on Obstetric Practice. Committee opinion no. 692: emergent therapy for acute-onset, severe hypertension during pregnancy and the postpartum period. *Obstet Gynecol.* 2017;129: e90–e95.

32. Tita AT, Szychowski JM, Boggess K, et al. Treatment for mild chronic hypertension during pregnancy. *N Engl J Med*. 2022;386:1781-1792.

33. Ormesher L, Higson S, Luckie M, et al. Postnatal enalapril to improve cardiovascular function following preterm preeclampsia (PICk-UP): a randomized double-blind placebo-controlled feasibility trial. *Hypertension*. 2020;76:1828–1837.

34. Smith RP. Diuretics and the treatment of preeclampsia. *Arch Intern Med.* 1982;142:1581.

35. Valensise H, Vasapollo B, Gagliardi G, Novelli GP. Early and late preeclampsia: two different maternal hemodynamic states in the latent phase of the disease. *Hypertension*. 2008;52:873-880.

36. Churchill D, Beevers GD, Meher S, Rhodes C. Diuretics for preventing pre-eclampsia. *Cochrane Database Syst Rev.* 2007;2007:CD004451.

37. Veena P, Perivela L, Raghavan SS. Furosemide in postpartum management of severe preeclampsia: a randomized controlled trial. *Hypertens Pregnancy*. 2017;36:84–89.

38. Ascarelli MH, Johnson V, McCreary H, Cushman J, May WL, Martin JN Jr. Postpartum preeclampsia management with furosemide: a randomized clinical trial. *Obstet Gynecol*. 2005;105:29-33.

39. Lopes Perdigao J, Lewey J, Hirshberg A, et al. Furosemide for accelerated recovery of blood pressure postpartum in women with a hypertensive disorder of pregnancy: a randomized controlled trial. *Hypertension*. 2021;77:1517-1524.

40. American College of Obstetricians and Gynecologists' Committee on Clinical Practice Guidelines—Obstetrics aKA, Gandhi M, Pettker CM, Simhan H. *Clinical Guidance for the Integration of the Findings of the Chronic Hypertension and Pregnancy (CHAP) Study.* 2022. Accessed November 27, 2023. https://www.acog.org/clinical/clinical-guidance/ for-the-integration-of-the-findings-of-the-chronichypertension-and-pregnancy-chap-study

41. Redman CW. Hypertension in pregnancy: the NICE guidelines. *Heart*. 2011;97:1967–1969.

42. Hauspurg A, Jeyabalan A. Postpartum preeclampsia or eclampsia: defining its place and management among the hypertensive disorders of pregnancy. *Am J Obstet Gynecol*. 2022;226:S1211-S1221.

43. Tucker KL, Mort S, Yu LM, et al. Effect of selfmonitoring of blood pressure on diagnosis of hypertension during higher-risk pregnancy: the BUMP 1 randomized clinical trial. *JAMA*. 2022;327: 1656-1665.

44. Chappell LC, Tucker KL, Galal U, et al. Effect of self-monitoring of blood pressure on blood pressure control in pregnant individuals with chronic or gestational hypertension: the BUMP 2 randomized clinical trial. *JAMA*. 2022;327:1666-1678.

45. Cairns AE, Tucker KL, Leeson P, et al. Selfmanagement of postnatal hypertension: the SNAP-HT trial. *Hypertension*. 2018;72:425-432.

46. Janssen MK, Demers S, Srinivas SK, et al. Implementation of a text-based postpartum blood pressure monitoring program at 3 different academic sites. *Am J Obstet Gynecol MFM*. 2021;3: 100446.

47. Hoppe KK, Williams M, Thomas N, et al. Telehealth with remote blood pressure monitoring for postpartum hypertension: a prospective single-cohort feasibility study. *Pregnancy Hypertens.* 2019;15:171–176.

48. van den Heuvel JFM, Lely AT, Huisman JJ, Trappenburg JCA, Franx A, Bekker MN. SAFE@-HOME: digital health platform facilitating a new care path for women at increased risk of preeclampsia - a case-control study. *Pregnancy Hypertens*. 2020;22:30–36.

49. Taylor RS, Freeman L, North RA. Evaluation of ambulatory and self-initiated blood pressure monitors by pregnant and postpartum women. *Hypertens Pregnancy*. 2001;20:25-33.

50. Kalafat E, Benlioglu C, Thilaganathan B, Khalil A. Home blood pressure monitoring in the antenatal and postpartum period: a systematic review meta-analysis. *Pregnancy Hypertens*. 2020;19:44–51.

51. Sanghavi M, Packard E, Sperling S, et al. Telemedicine may increase visit completion rates in postpartum patients with preeclampsia. *PLoS One*. 2022;17:e0275741.

52. Payakachat N, Rhoads S, McCoy H, Dajani N, Eswaran H, Lowery C. Using mHealth in post-

partum women with pre-eclampsia: lessons learned from a qualitative study. *Int J Gynaecol Obstet*. 2020;149:339-346.

53. de Mooij MJM, Hodny RL, O'Neil DA, et al. OB nest: reimagining low-risk prenatal care. *Mayo Clin Proc.* 2018;93:458-466.

54. Rivera-Romero O, Olmo A, Munoz R, Stiefel P, Miranda ML, Beltran LM. Mobile health solutions for hypertensive disorders in pregnancy: scoping literature review. *JMIR Mhealth Uhealth.* 2018;6: e130.

55. Hirshberg A, Downes K, Srinivas S. Comparing standard office-based follow-up with text-based remote monitoring in the management of post-partum hypertension: a randomised clinical trial. *BMJ Qual Saf.* 2018;27:871-877.

56. Hirshberg A, Sammel MD, Srinivas SK. Text message remote monitoring reduced racial disparities in postpartum blood pressure ascertainment. *Am J Obstet Gynecol.* 2019;221:283–285.

57. Khosla K, Suresh S, Mueller A, et al. Elimination of racial disparities in postpartum hypertension follow-up after incorporation of telehealth into a quality bundle. *Am J Obstet Gynecol MFM.* 2022;4:100580.

58. Rossiter C, Henry A, Roberts L, et al. Optimising mothers' health behaviour after hypertensive disorders of pregnancy: a qualitative study of a postnatal intervention. *BMC Publ Health.* 2022;22:1259.

59. Rich-Edwards JW, Stuart JJ, Skurnik G, et al. Randomized trial to reduce cardiovascular risk in women with recent preeclampsia. *J Womens Health (Larchmt).* 2019;28:1493–1504.

60. Herring SJ, Bersani VM, Santoro C, McNeil SJ, Kilby LM, Bailer B. Feasibility of using a peer coach to deliver a behavioral intervention for promoting postpartum weight loss in Black and Latina mothers. *Transl Behav Med*. 2021;11:1226-1234.

61. Center for Disease Control Policy Evidence Assessment Report: Community Health Worker Policy Components. 2014. Accessed November 27, 2023. https://www.cdc.gov/dhdsp/pubs/ docs/chw_evidence_assessment_report.pdf

62. Bellad MB, Goudar SS, Mallapur AA, et al. Community level interventions for pre-eclampsia (CLIP) in India: a cluster randomised controlled trial. *Pregnancy Hypertens*. 2020;21:166–175.

63. Bone JN, Magee LA, Singer J, et al. Blood pressure thresholds in pregnancy for identifying

maternal and infant risk: a secondary analysis of Community-Level Interventions for Pre-eclampsia (CLIP) trial data. *Lancet Glob Health*. 2021;9: e1119-e1128.

64. McCue K, Sabo S, Wightman P, et al. Impact of a community health worker (CHW) home visiting intervention on any and adequate prenatal care among ethno-racially diverse pregnant women of the US Southwest. *Matern Child Health J.* 2022;26: 2485–2495.

65. Barsky J, Hunter R, McAllister C, et al. Analysis of the implementation, user perspectives, and feedback from a mobile health intervention for individuals living with hypertension (DREAM-GLOBAL): mixed methods study. *JMIR Mhealth Uhealth.* 2019;7:e12639.

66. Jack HE, Arabadjis SD, Sun L, Sullivan EE, Phillips RS. Impact of community health workers on use of healthcare services in the United States: a systematic review. *J Gen Intern Med.* 2017;32: 325–344.

67. Lewin S, Munabi-Babigumira S, Glenton C, et al. Lay health workers in primary and community health care for maternal and child health and the management of infectious diseases. *Cochrane Database Syst Rev.* 2010;2010:CD004015.

68. Kiely B, Croke A, O'Shea M, et al. Effect of social prescribing link workers on health outcomes and costs for adults in primary care and community settings: a systematic review. *BMJ Open.* 2022;12:e062951.

69. Integrating Social Care into the Delivery of Health Care: Moving Upstream to Improve the Nation's Health. 2019. Accessed November 27, 2023. https://nap.nationalacademies.org/catalog/25467/ integrating-social-care-into-the-delivery-of-healthcare-moving

70. Hameed AB, Haddock A, Wolfe DS, et al. Alliance for innovation on maternal health: consensus bundle on cardiac conditions in obstetric care. *Obstet Gynecol*. 2023;141:253-263.

71. Suresh SC, Duncan C, Kaur H, et al. Postpartum outcomes with systematic treatment and management of postpartum hypertension. *Obstet Gynecol.* 2021;138:777-787.

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