

Pregnancy-Related Optimal Management of Hypertension (PROMPT)

The PROMPT initiative's purpose is to assist Florida maternity hospitals and providers improve maternal-health outcomes by implementing standardized evidence-based practices for the identification, management, and treatment of severe hypertension (HTN) in pregnancy and postpartum across healthcare settings.

Hospitals will report:

- 1. Aggregate Severe Maternal Morbidity (SMM) overall as well as among preeclampsia, eclampsia, and HELLP syndrome cases.
- 2. Abstracted patient data:
 - The first 10 cases* of pregnant and postpartum patients that present to EMS/ED, OBED, antepartum, L&D, or postpartum unit with Persistent Severe Hypertension.
 - The first 5 cases* of pregnant and postpartum patients during their delivery admission with preeclampsia, eclampsia, or HELLP syndrome who experienced SMM (exclude transfusion-only cases).
 - *NOTE: hospitals will submit the number of cases specified or as many as they have in the month.

 Cases included in the second group (SMM) must be excluded from the first to avoid submitting duplicate patients.
- **3. Hospital-level measures** (policies, procedures, or guidelines to increase hospital's capacity to implement the initiative and staff education and training).

The measures listed in this document will be calculated and reported monthly to participating hospitals in a quality improvement data report so that facilities can track their progress. These measures are subject to change during the process of finalizing data collection and reporting tools.

#	OUTCOME MEASURES	Description	Reported	Source
01	Severe Maternal Morbidity (excluding transfusion codes alone)	Denominator : patients during their delivery admission. Numerator : among the denominator, those who experienced severe maternal morbidity, excluding those who experienced transfusion alone. Find SMM definition and ICD-10 codes here .	Quarterly	Abstracted from medical chart
O2	Severe Maternal Morbidity among patients with preeclampsia, eclampsia, and HELLP syndrome (excluding transfusion codes alone)	Denominator: patients during their delivery admission with preeclampsia, eclampsia, or HELLP syndrome (ICD-10 codes). Numerator: among the denominator, those who experienced severe maternal morbidity (SMM), excluding those who experienced transfusion alone. Find SMM definition and ICD-10 codes here.	Quarterly	Abstracted from medical chart
#	PROCESS MEASURES	Description	Reported	Source
P1	Timely treatment of persistent severe hypertension Two measures for timely treatment will be reported: 1. % Patients identified and treated in EMS/ED, and 2. % Patients identified and treated in an OB unit (OBED, AP, L&D, PP)	Denominator: pregnant and postpartum patients with persistent severe hypertension. * Numerator: among the denominator, those who were treated within 1 hour with IV Labetalol, IV Hydralazine, PO Nifedipine, or an antihypertensive approved in your unit protocol. Note: The "1 hour" is measured from the first severe-range BP reading. *Persistent Severe Hypertension is defined as BP ≥160 systolic and/or ≥110 diastolic with: 1. one or more repeat severe HTN observations 15-60 minutes after onset (values do not need to be consecutive), or 2. BP not documented to have decreased to non-severe HTN within 15 minutes.	Monthly – Patient sample Disaggregated by race-ethnicity, insurance type, language and age Average time to treatment will be reported as an adjunct measure Full measurement specifications: SMFM Special Statement	Abstracted from medical chart Refer to FPQC FAQs for additional guidance

#	PROCESS MEASURES	Description	Reported	Source
P2	Antihypertensive administration per unit protocol	Denominator: # qualifying patients in sample. Numerator: among the denominator, those who received an antihypertensive medication per unit protocol (dosage and timing). Note: Your unit should have a protocol in place for antihypertensive medication administration. This protocol should have been reviewed within the past year and align with current ACOG recommendations.	Monthly – Patient sample	Abstracted from medical chart
P3	Physician and nurse debrief	Denominator : # qualifying patients in sample. Numerator : among the denominator, the number of cases where the physician and nurse conducted a debrief after the severe event. See AWHONN debrief form here.	Monthly – Patient sample	Abstracted from medical chart
P4	Interdisciplinary review of SMM cases	Denominator: # of patients who experienced a Severe Maternal Morbidity (SMM) event. Numerator: Number of SMM cases for which a referral was made for interdisciplinary case review. Note: Referrals are typically made to the hospital's Quality Assurance (QA) or Patient Safety Committee, Maternal Mortality or Morbidity Review Committee, or a designated Interdisciplinary Review Team responsible for case evaluations and system-level learning.	Monthly – Patient sample	Abstracted from medical chart
P5	*Measure combines P6 and P7 below	Denominator: # qualifying patients in sample Numerator: among the denominator, those who received 1. Verbal and written education on Urgent Postpartum Warning Signs; and 2. Patient verbal debrief by physician and nurse.* *Patient debrief definition: a structured conversation where the physician and nurse share with the patient the details of SHTN event, including any complications, treatment, and follow-up care.	Monthly – Patient sample	Abstracted from medical chart

#	PROCESS MEASURES	Description	Reported	Source
P6	Patient given verbal and written education on urgent postpartum warning signs	Denominator : # qualifying patients in sample. Numerator : among the denominator, those who received verbal and written Urgent Postpartum Warning Signs.	Monthly – Patient sample	Abstracted from medical chart
P7	Patient verbally debriefed by physician and nurse	Denominator: # qualifying patients in sample. Numerator: among the denominator, those who were verbally debriefed* by physician and nurse. *Patient debrief definition: a structured conversation where the physician and nurse share with the patient the details of SHTN event, including any complications, treatment, and follow-up care. "• Include patient support networks during patient event debriefs, as requested. • This measure is not intended to represent a disclosure conversation but rather reflects a standard part of care that is a discussion between the patient and their care team." (AIM, 2024)	Monthly – Patient sample	Abstracted from medical chart
P8	BP cuff provision or validation prior to discharge	Denominator: # qualifying patients in sample. Numerator: among the denominator, the number of patients who either: 1. Received a BP cuff to take home, or 2. Brought a BP cuff in from home to be validated.	Monthly – Patient sample	Abstracted from medical chart
P9	Scheduled follow-up within 7 days after discharge	Denominator: # qualifying patients in sample. Numerator: among the denominator, those with a scheduled follow-up, including blood pressure and symptoms check, within 7 days after their discharge date. Includes patients with weekend discharges. Excludes patients transferred out before discharge. Note: Two separate periods will be tracked: 1. % patients with BP and symptom check within 3 days, and 2. % patients with scheduled follow-up 4-7 days post discharge.	Monthly – Patient sample	Abstracted from medical chart

#	PROCESS MEASURES	Description	Reported	Source
P10	Provider and nurse education using initiative, promotional and educational materials	Training bundle includes education on: 1. accurate blood pressure measurement & assessment; 2. severe hypertension/preeclampsia policy, guidelines or procedures; 3. respectful care and commitment to respectful-care practices. Report separately the percentage of physicians/midwives and nurses who have completed education on each topic of the training bundle since the PROMPT kickoff.	Quarterly	Varies per hospital (tally, system report, etc.) Hospital-level data
#	BALANCING MEASURES	Description	Reported	Source
B1	Hypotension and hypotensive- related events following antihypertensive medication	Percentage of patients who developed hypotension after antihypertensive administration for persistent SHTN. Denominator: # patients given antihypertensive medication for persistent SHTN. Numerator: among the denominator, patients with documented hypotension within one hour of antihypertensive administration for persistent SHTN. Among those that develop hypotension after antihypertensive medication for persistent SHTN, the following additional measures will be reported: - Fetal heart rate deterioration related to hypotension - Clinical intervention required for hypotensive episodes - Cesarean deliveries performed due to hypotension events	Monthly – Patient sample	Abstracted from medical chart

The following measures are collected to provide additional context about adherence to medication protocols and clinical decision-making in managing severe hypertension. Because policies, processes, or guidelines may vary across units, please refer to your unit-specific protocols when assessing adherence.

- 1. **Use of Specific Antihypertensive Medications -** *Labetalol, Hydralazine, Nifedipine, other antihypertensives listed in your unit protocol.*
- 2. **First Antihypertensive Medication Administered** *Identifies the initial medication choice for persistent SHTN management.*
- 3. **Use of Magnesium Sulfate** The measure will reflect the **percentage of all patients in the sample** who received magnesium sulfate, regardless of individual clinical indication. While not all patients require magnesium sulfate—its use depends on physician clinical judgment—tracking it across the full sample supports consistent, population-level monitoring of practice patterns and helps identify variation or potential gaps in care across sites.
- 4. **Use of Magnesium Sulfate for Patients with Preeclampsia, Superimposed Preeclampsia, Eclampsia, and/or HELLP** The measure will reflect the percentage of patients with Preeclampsia, Superimposed Preeclampsia, Eclampsia, and/or HELLP that received magnesium sulfate. While not all

<u>patients require magnesium sulfate</u>—its use depends on physician clinical judgment—tracking it across the full sample supports consistent, population-level monitoring of practice patterns and helps identify variation or potential gaps in care across sites.

- 5. **Magnesium Sulfate Administration per Unit Protocol** *Follow your unit protocol to determine if protocol was met.*
- 6. **Clinical Reasons for Not Administering Antihypertensive/Magnesium Sulfate** Reasons including: clinical judgment, patient preference, BP improvements, or other clinical decisions.

STRUCTURAL MEASURES

Hospitals need to implement and/or reinforce key processes, guidelines, policies, and resources to support PROMPT. Hospitals will report structural measures until they have them fully implemented. Quarterly updates are required. Report as follows:

- Not Started
- Planning
- Started Implementing started implementation in the last 3 months
- Implemented less than 80% compliance after at least 3 months of implementation (not routine practice)
- Fully Implemented at least 80% compliance after at least 3 months of implementation (routine practice)

#	STRUCTURAL MEASURES	Description	Source
S1	Emergency Department (ED) screening for current or recent pregnancy	ED-established standardized verbal screening for pregnancy now and during the past year as part of its triage or initial assessment process. Note: This verbal screening recommendation is based on timing, not the outcome of the pregnancy. Details on the pregnancy outcome are not required.	Quarterly Hospital-level data
S2	Implement a process to ensure accurate blood pressure measurement and assessment with confirmation after severe range	Implement a process to ensure staff is educated on how to accurately measure and assess blood pressure with a confirmation after severe range.	Quarterly Hospital-level data
S3	Implement a standard protocol, guidelines and/or processes for identification, management, and treatment of severe HTN	 Implementation of a Severe Hypertension/Preeclampsia policy, guidelines or procedure (reviewed and updated in the last 2 years) that contain the following: Treatment of SHTN/preeclampsia The use of seizure prophylaxis, including for the treatment of magnesium overdose 	Quarterly Hospital-level data
S4	Ready reference algorithm	Ensure ready reference to algorithms for identifying, assessing, and treatment SHTN/preeclampsia on all units.	Quarterly Hospital-level data
S5	Level of care escalation	Implemented a system plan for level of care escalation, consultation, and maternal transport when needed.	Quarterly Hospital-level data

S6	Rapid access to medication	Develop a workflow to ensure rapid access to SHTN medication.	Quarterly Hospital-level data
S7	Establish a standardized process to conduct debriefs with patients after a severe event	 AIM recommendations: Include patient support networks during patient-event debriefs, as requested. This measure is not intended to represent a disclosure conversation but rather reflects a standard part of care that is a discussion between the patient and the care team. Severe events may include The Joint Commission sentinel event definition, severe maternal morbidity, or fetal death. 	Quarterly Hospital-level data
S8	Interdisciplinary case reviews	Establish a process to perform interdisciplinary systems-level reviews of cases of severe maternal morbidity (including, at a minimum, pregnant and postpartum patients admitted to the ICU or who received ≥ 4 units RBC transfusions).	Quarterly Hospital-level data
S9	Drills and simulations	Hold interdisciplinary and interdepartmental team-based drills with timely debriefs that include the use of simulated patients. Severe Hypertension Drills should cover all sequelae, such as preeclampsia.	Quarterly Hospital-level data
S10	Implement periodic education and engagement for ED physicians and staff about Severe Hypertension/Preeclampsia	Develop a strategy to engage and educate ED physicians & staff on signs and symptoms of severe hypertension and preeclampsia in pregnant and postpartum patients.	Quarterly Hospital-level data
S11	Engage a patient advisor in the QI team	Identify, engage, and onboard a patient advisor in your QI team. Provide a clear role description, goals, contact information, whether it will be an inperson/virtual/hybrid role, and a meeting schedule. Patient advisors may assist by sharing insights from lived experiences, reviewing materials, enhancing trust and communication with patients, helping to align care strategies with patient needs, promoting shared decision-making, identifying care gaps from the patient's perspective, etc. Find resources in the FPQC toolbox.	Quarterly Hospital-level data

Questions? Please contact FPQC@usf.edu v. 5/1/2025