



Preeclampsia in the ED: Rapid Recognition, Timely Response

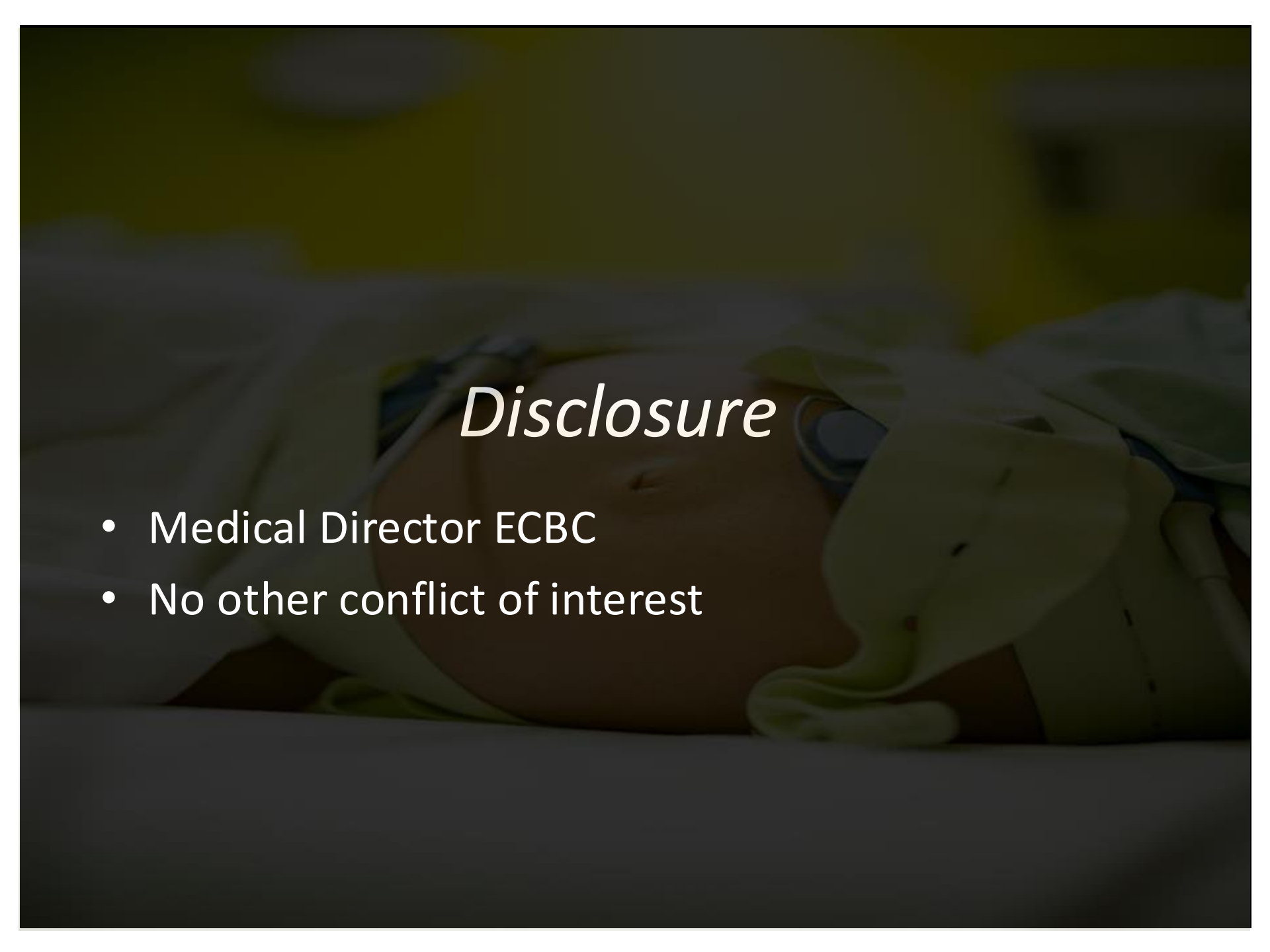
St Paul's Emergency Medicine Update
Presented by: Dr. Todd Ring
Sept 2025

Presenting today's topic from the unceded territories of the Lílwat Nation and the Skwxwú7mesh Úxwumixw

For our Team at Emergency Care BC, acknowledging that we are on the traditional territories of First Nations communities across our province is an expression of cultural humility – and an understanding that we are privileged to use and share this land – in addition to recognizing our duty and desire to provide culturally safe care to First Nations, Inuit, and Métis people across BC.



EMERGENCY CARE BC
Provincial Health Services Authority

A newborn baby is lying in a hospital bed, partially covered by a white blanket. The baby's face is visible, and they appear to be sleeping. Medical equipment, including a nasal cannula and a pulse oximeter, is attached to the baby. The background is a plain, light-colored wall.

Disclosure

- Medical Director ECBC
- No other conflict of interest

Disclaimer

In alignment with PHSA fiscal responsibilities regarding travel, no PHSA funding was used to support ECBC participation at St. Paul's Emergency Medicine conference.

ECBC remains committed to improving emergency medicine care provincially and looks forward to increased in-person presence at future events.



emergencycarebc.ca/spemu2025

Notice of Recording

This session is being recorded
to explore AI's role in increasing the efficiency of knowledge translation.

*Click the QR to access session tools and resources.
Watch for a summary of this session supported by AI coming soon.*



emergencycarebc.ca/spemu2025

Objectives

Recognize

Recognize clinical presentation and diagnostic criteria of preeclampsia

Initiate

Initiate ED-based management and stabilization of preeclampsia/eclampsia and severe HTN

Apply

Apply Canadian guidelines in ED workflows

Main Take-Aways

- Clear antenatal emergency care pathways
 - ED/L&D
- Consider preeclampsia in any pregnant/immediate post partum patient
 - Elevated BP (SBP ≥ 140 or DBP ≥ 90) and *any* end organ dysfunction
- Early BP management to prevent complications
 - ED aggressive treatment of severe HTN (SBP ≥ 160 or DBP ≥ 110)
- MgSO₄ 4gm IV eclampsia prophylaxis/treatment





Clinical Case: ED Presentation

28-year-old G1P0 at 32 weeks gestation presents with:

- Severe headache and visual disturbances
- BP: 165/100 mmHg
- HR: 92 bpm, afebrile
- Mild right upper quadrant tenderness
- No prior diagnosis of hypertension

**Triage nurse asks:
Should we triage patient to
ED or to L&D?**

Questions

1. Where should this patient be seen?
2. What medication should this patient immediately receive?
3. You are working in a rural facility without OB. What should you consider pre-transport?

Three Vessels and Trachea View

Four-Chamber View

Five-Chamber View

Left Ventricular Short Axis View

Short Axis of the Great Vessels / RVOT

Abdomen View

CRANIAL

CAUDAL

Ductal Arch

Aortic Arch

Venae Cavae

Hypertension

- Hypertension SBP \geq 140 or DBP \geq 90
 - Transient/white coat and masked
 - Severe HTN SBP \geq 160 or DBP \geq 110
- Proteinuria
 - Definitive testing: random urine PCR; ACR or 24h protein
 - Urine dipstick: \geq 2+ protein

2020
HYPERTENSION
HIGHLIGHTS



A Practical Guide informed
by the Hypertension Canada
Guidelines for the Prevention,
Diagnosis, Risk Assessment,
and Treatment of Hypertension



Hypertensive Disorders of Pregnancy



Chronic HTN

Preexisting HTN or BP > 140/90 mm Hg before 20 wks of gestation, or persist > 12 wks post partum



Gestational HTN

New HTN (BP > 140/90 mm Hg) after 20 wks of gestation
No end organ dysfunction



Preeclampsia

Gestational HTN + Proteinuria.
Low platelets, High LFTs, Pulm edema, visual disturbances

Eclampsia

Preeclampsia + seizures during pregnancy or within 10 days post partum



43047-10-07-26-15

RA-4-8UOB

M 1.2 Clinica "STAIKA" Cluj Napoca

13.6cm / 5.37

Tls 0.3

10-12-2010 11:47:30

Pwr 100 %
Gn -2
WMF 190 Hz
SV Angle 0
Size 3.0mm
Frq low
PRF 5.5kHz



Umb-PS	30.45cm/s
Umb-ED	-7.77cm/s
Umb-S/D	-3.92
Umb-PI	3.29
Umb-RI	1.26
Umb-MD	-7.77cm/s
Umb-I Amax	11.62cm/s
Umb-IR	130bpm

Diagnostic Criteria

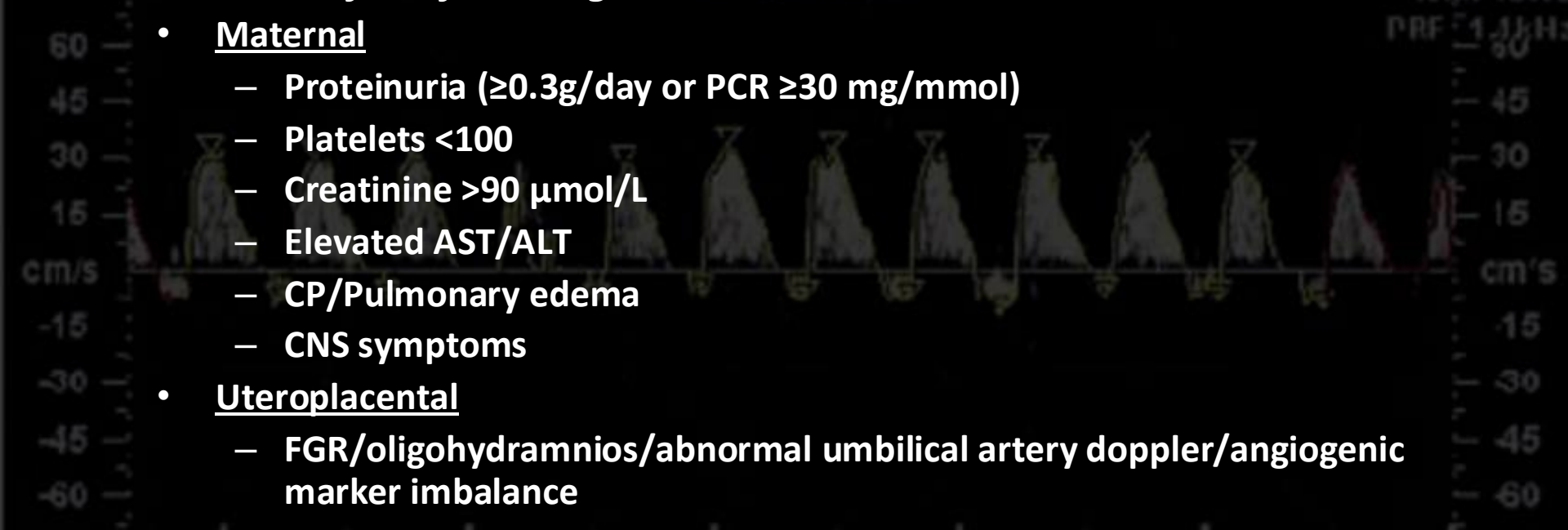
- BP $\geq 140/90$ mmHg on 2 occasions, 4h apart
AND one of the following:

- Maternal

- Proteinuria ($\geq 0.3g/day$ or PCR ≥ 30 mg/mmol)
- Platelets < 100
- Creatinine > 90 $\mu\text{mol/L}$
- Elevated AST/ALT
- CP/Pulmonary edema
- CNS symptoms

- Uteroplacental

- FGR/oligohydramnios/abnormal umbilical artery doppler/angiogenic marker imbalance

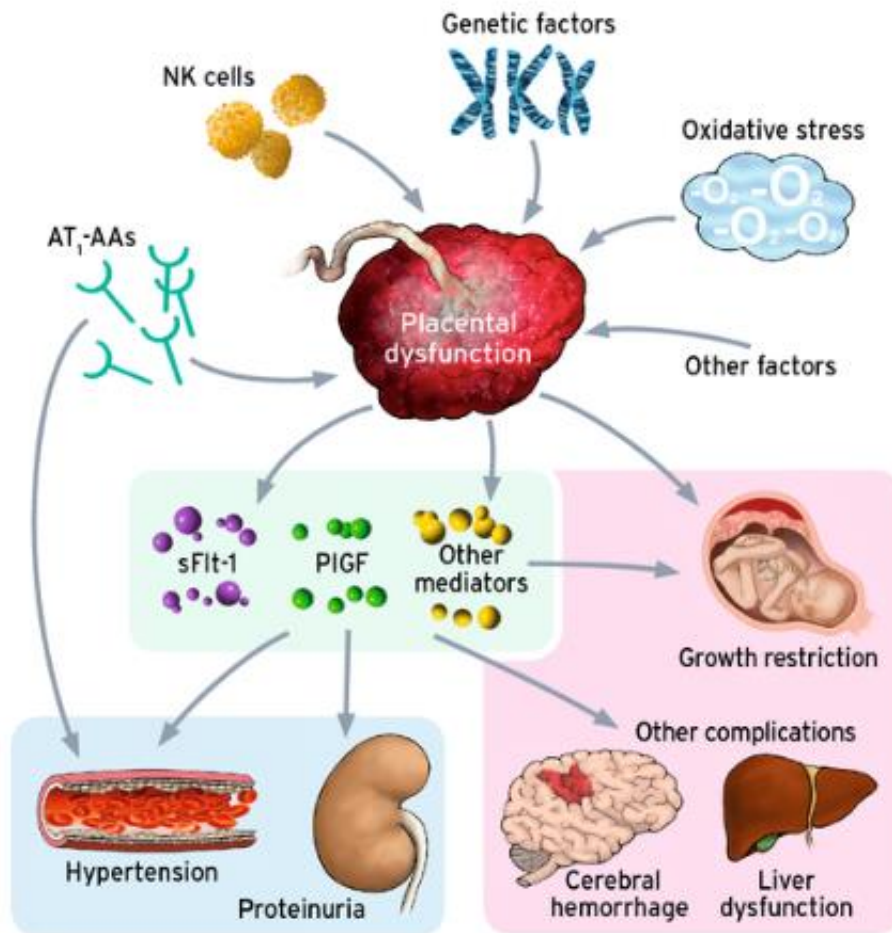




Why It Matters

- Preeclampsia affects ~5% of pregnancies
- Significant contributor to maternal morbidity and mortality
- EDs are critical access points for undiagnosed or worsening cases

Pathophysiology



- **Abnormal placentation**
- **Placental under-perfusion/ ischemia**
- **Maternal endothelial dysfunction**
- **Complications:**
 - **Maternal: HTN, vasospasm, platelet aggregation**
 - **Fetal: hypoperfusion**

When to Suspect Preeclampsia?







- Pregnant ≥ 20 weeks or up to 6 weeks postpartum with:
 - New or worsening hypertension
 - Headache, vision changes, RUQ pain
 - Dyspnea, edema, seizure, altered LOC









Risk Factors

CLINICAL RISK FACTORS FOR PREECLAMPSIA (EARLY PREGNANCY)

High-risk factors (any 1)

-  Prior preeclampsia
-  Pre-pregnancy BMI $>30 \text{ kg/m}^2$
-  Chronic hypertension
-  Pre-gestational diabetes mellitus
-  Chronic kidney disease
-  Systemic lupus erythematosus /antiphospholipid antibody syndrome

Moderate-risk factors (2 needed)

-  Prior placental abruption
-  Prior stillbirth
-  Prior fetal growth restriction (FGR)
-  Maternal age $>40 \text{ y}$
-  Nulliparity
-  Multifetal pregnancy

Women are at increased risk if they have ≥ 1 high-risk factor or ≥ 2 moderate-risk factors.

(Adapted from Magee et al., ISSHP 2021 Guidelines)


A pregnant woman is shown in profile, sitting up in a hospital bed. She is wearing a light-colored hospital gown. The background is dimly lit, showing medical equipment and monitors. The overall tone is serious and clinical.

**PRIORITIES OF
ED MANAGEMENT OF
PREECLAMPSIA/
ECLAMPSIA**

ED Management Priorities



- **Clear ED/L&D triage guideline**
- **ABCs – stabilize first**
- **BP control**
- **Seizure prophylaxis**
- **OB consult and admission**
 - **Delivery definitive cure**
- **Transportation consideration**



PERINATAL SERVICES BC REVIEW: ED/L&D GUIDELINES



Perinatal Triage Guidelines



Perinatal Patients Presenting to the Emergency Department - Royal Inland Hospital

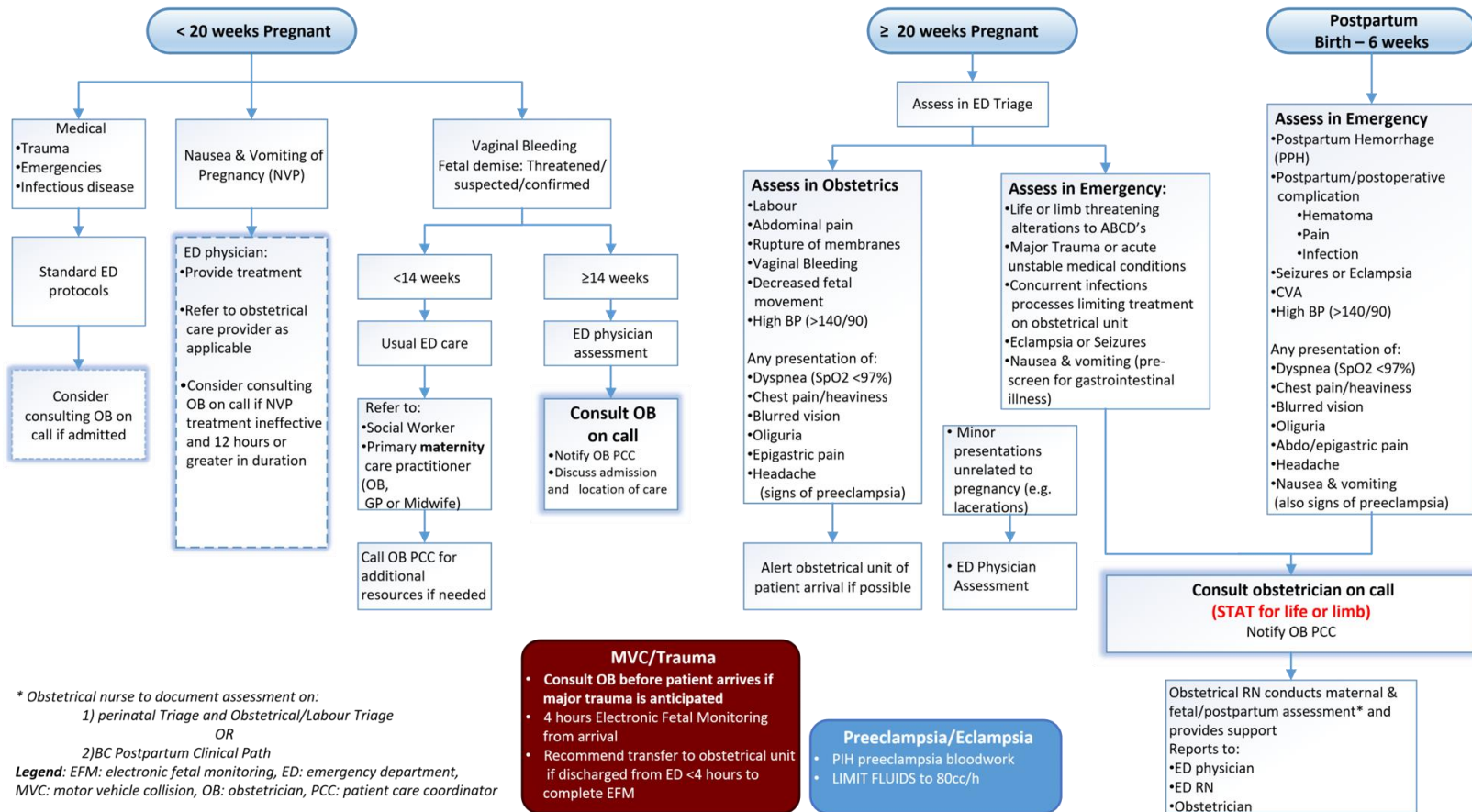


Figure 2. Suggested dose titration of antihypertensive therapy for urgent control of hypertension in pregnancy^a.

Drug	Caution	T 0 min	T 30 min	T 60 min	T 90 min	T 120 min	T 150 min	T 180 min
Labetalol (oral)	• Contraindicated in patients with uncontrolled asthma or heart failure	200 mg	—	200 mg	—	200mg	—	
Labetalol (IV intermittent)	• Caution with hypoglycemic unawareness in diabetes	10–20 mg	20–40 mg ^b	40–80 mg	40–80 mg	40–80 mg	40–80 mg ^c	
Labetalol (IV infusion)	• May cause neonatal bradycardia and neonatal hypoglycemia and warrants newborn screening	0.5–2 mg/min	→	→	→	→	→ ^d	
Nifedipine (oral capsule swallowed whole, not bitten or crushed)	• May cause maternal headache and tachycardia	5-10 mg	10 mg	—	10 mg	—	10 mg	
Methyldopa (oral)	• Onset of action may be delayed	1000 mg	—	—	—	—	—	
Hydralazine (IV)	• May increase risk of maternal hypotension, and maternal and fetal tachycardia	5 mg	5–10 mg	5–10 mg ^e	5–10 mg ^e			

Use alternative from a different drug class^d

Reproduced with permission from Magee LA, Brown MA, Hall DR, et al. The 2021 International Society for the Study of Hypertension in Pregnancy classification, diagnosis & management recommendations for international practice. *Pregnancy Hypertens.* 2021;27:148-69.¹³⁴

^a When severe hypertension has resolved, switch to routine oral medication.

^b Double the initial dose of labetalol IV.

^c Do not exceed the maximum dose of IV labetalol, which is 300 mg total in a treatment course.

^d If nifedipine or hydralazine were the initial drug used, choose oral labetalol or oral methyldopa as the alternative.

^e Do not exceed the maximum dose of IV hydralazine of 20 mg.

IV: intravenous.

Figure 1. Maintenance therapy and suggested dose titration of antihypertensive therapy for non-urgent control of hypertension in pregnancy.

First-line drug	Caution	Low ^a	Dosage (mg)				
			If BP not controlled	Medium	If BP not controlled on medium dosage	High ^b	Maximum
Labetalol	<ul style="list-style-type: none"> Contraindicated with poorly controlled asthma Caution with hypoglycemic unawareness in diabetes May cause neonatal bradycardia and hypoglycemia and warrants new born screening 	100 TID or QID	proceed to medium dose of same low-dose medication	200 TID or QID	Consider adding another low-dose medication rather than going to a high-dose of the same medication(s), for a maximum of 3 medications	300 TID or QID	1200/d
Nifedipine XL	<ul style="list-style-type: none"> Contraindicated with aortic stenosis Ensure extended release (XL) formulation 	30 OD	proceed to medium dose of same low-dose medication	30 BID or 60 OD	Consider adding another low-dose medication rather than going to a high-dose of the same medication(s), for a maximum of 3 medications	30 QAM and 60 QPM	120/d
Methyldopa	<ul style="list-style-type: none"> May cause maternal depression 	250 TID or QID	proceed to medium dose of same low-dose medication	500 TID-QID	Consider adding another low-dose medication rather than going to a high-dose of the same medication(s), for a maximum of 3 medications	750 TID	2500/d

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^a Starting doses are higher than generally recommended for adults, given more rapid clearance in pregnancy.

^b When a medication is at high (or maximum) dosage, consider using a different medication to treat any severe hypertension that may develop.

Source: adapted from ALARM 27th Edition ALARM Manual, Table 8 of the SOGC 2014 guideline and Magee et al.¹³⁶

OD: once daily; TID: 3 times daily; QAM: every morning; QID: 4 times daily; QPM: every evening.

Figure 3. Magnesium sulphate dosage and monitoring.

Dosage ^{35, 136}	IV administration	Combined IV and IM administration ^a
Loading dose	4 g MgSO ₄ IV in 100 mL normal saline solution, infused over 20 min using an infusion device	4 g MgSO ₄ IV in 100 mL normal saline solution, infused over 20 min using an infusion device <i>and</i> 5 g IM into <i>each</i> buttock (for a total of 10 g), every 4 h
Maintenance	1 g/h IV in normal saline solution, using an infusion device	5 g IM into <i>one</i> buttock every 4 h
Duration	Until 24 h after last eclamptic seizure or birth, whichever is later	
Monitoring	Observations	Signs of toxicity ^b
Maternal		
Upon completion of loading dose	Reflexes	Decreased or absent
Every 30 min	BP	Lower
	Respiratory rate	Lower or cardiac arrhythmias
	Pulse oximetry	<12/min for 15 min
Every 4 h	Urine output ^d	O ₂ saturation <94% for 15 min
	Reflexes	<30 mL/h for 4 h ^e
Symptoms ^f	Central nervous system (e.g., excessive drowsiness, slurred speech)	Decreased or absent
	Neuromuscular (e.g., muscle weakness)	
Fetal		
≥26 wk	Continuous cardiotocography	
<26 wk	Intermittent FHR auscultation every 30 min	

MgSO₄ 4gms IV then

1gm/h

Transport:

MgSO₄ 4gm IV and

5gm IM each buttock

followed by 5gm IM

one buttock q4h

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^a Administration can be switched to IV dosing by starting 1 g/h (without a loading dose) when the next dose of IM magnesium sulphate is due.

^b Monitoring of serum magnesium levels is not necessary unless there is decreased renal function or signs of toxicity.

^c If toxicity is suspected, cease the MgSO₄ infusion and take blood for serum Mg level. If toxicity is clear, administer calcium gluconate 10% (10 mL in 100 mL normal saline solution IV over 3 min).

^d Foley catheterization is recommended.

^e Decreased urine output is included because it increases the risk of toxicity.

^f Symptoms of toxicity should be distinguished from well-known side effects, which include flushing of the skin, a metallic taste in the mouth, sweating, nausea and vomiting, heaviness in the chest, palpitations, and lowering of the BP initially.

BP: blood pressure; FHR: fetal heart rate; MgSO₄: magnesium sulphate; O₂: oxygen.

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BP: blood pressure; FHR: fetal heart rate; MgSO₄: magnesium sulphate; O₂: oxygen.

Treatment of Preeclampsia

- **Delivery only cure**
- **SOGC no longer defines as “severe preeclampsia”**
- **Need to consider maternal and fetal well being and gestational age in decision to deliver**



Timing of Delivery

	Viability to 33 ⁶ weeks	34 ⁰ — 36 ⁶ weeks	≥ 37 ⁰ weeks
Chronic HTN	Expectant	Expectant	Offer at 38 ⁰ ; advise at 40 ⁰
Gestational HTN	Expectant	Expectant	If HTN arose at <37 ⁰ <ul style="list-style-type: none"> • offer at 38⁰; • advise at 40⁰ If HTN arises at ≥37 ⁰ initiate delivery
Preeclampsia	Expectant	34 ⁰ — 35 ⁶ Discuss (maternal benefit vs. fetal risk) 36 ⁰ — 36 ⁶ Initiate delivery	Initiate delivery

Adopted from Table 6: SOGC Guideline #26 2022

Workup in the ED

- Labs: CBC, electrolytes, creatinine, LFTs, urinalysis
- Urine protein/creatinine ratio
- ECG, chest X-ray if respiratory or cardiac symptoms
- Head CT if neurologic symptoms or seizures

Disposition from the ED

- All suspected cases
→ OB consult +
admission
- Stabilize and arrange
transfer if OB not
onsite
- Protocols for
rural/remote settings
- Consider fetal
monitoring if viable
gestation



emergency exit

Transport

Transport Checklist for Women with Preeclampsia (Condensed)

Factor	Key Actions
Maternal stabilized	BP <160/110, antihypertensives given, IV access consider Foley, patient responsive/intubated if required
Fetal status	Document FHR (present/absent); if present, no delivery indication for transport
Eclampsia prophylaxis	MgSO ₄ (5 g IM each buttock or IV), Calcium gluconate available
Skilled provider	Monitor BP & reflexes hourly, manage seizures (MgSO ₄ , airway, ventilation), monitor SpO ₂ , document FHR before/after transport
Meds available	Nifedipine 5 mg cap / Labetalol 200 mg tab / MgSO ₄ .4 g IV
Confirm with receiving centre	Tocolysis if needed, antenatal steroids if <34-6 weeks



Special Considerations

- **Postpartum preeclampsia: up to 6 weeks postpartum**
- **Eclampsia: seizures may be first sign**
- **HELLP syndrome can occur without hypertension**
- **Magnesium + BP meds should be available in all EDs**

Clinical Case: ED Presentation

- **28-year-old G1P0 at 32 weeks gestation presents with:**

- Severe headache and visual disturbances
- BP: 165/100 mmHg
- HR: 92 bpm, afebrile
- Mild right upper quadrant tenderness
- No prior diagnosis of hypertension

Triage nurse asks: Should we triage patient to ED or to L&D?

Three Vessels and Trachea View

Four-Chamber View

Five-Chamber View

Left Ventricular Outflow Tract

Short Axis of the Great Vessels / RVOT

Abdomen View

CROWIAL

CAUDAL

Ductal Arch

Aortic Arch

Venae Cavae

Questions

1. Where should this patient be seen?

- a. Emergency department
- b. Labor and delivery
- c. Who cares just call OB

Three Vessels and Trachea View

Four-Chamber View

Five-Chamber View

Left Ventricular Outflow Tract

Short Axis of the Great Vessels / RVOT

Abdomen View

Cranial

Caudal

Ductal Arch

Aortic Arch

Venae Cavae

Questions

2. What medication should this patient immediately receive?

– MgSO₄ 4 gm IV followed by 1gm IV/h

or

– MgSO₄ 4gm IV followed by 5gm IM each buttock

and

– Labetolol 10 – 20 mg IV or Labetolol 200mg po

Questions

3. You are working in a rural facility without OB. What should you consider pre-transport?

- a. Maternal stabilized
- b. Fetus stabilized
- c. Prophylaxis given
- d. Provider
- e. Meds available
- f. Receiving center

Three Vessels and Trachea View

Four-Chamber View

Five-Chamber View

CROWIAL

Axis of the Great Vessels / RVOT

Abdomen View

CAUDAL

Ductal Arch

Aortic Arch

Venae Cavae

5 TAKE HOME SUMMARY POINTS

- 1. Think preeclampsia in any pregnant/postpartum patient with systemic symptoms**
- 2. Use BP + one organ dysfunction for diagnosis**
- 3. Start magnesium and antihypertensives early**
- 4. Communicate with OB and arrange timely disposition**
- 5. Implement ED-specific protocols**

Questions?



ED QUICK REFERENCE CHECKLIST



Suspect in pregnant ≥ 20 weeks or ≤ 6 weeks postpartum with headache, vision changes, RUQ pain, dyspnea, edema, seizures



BP $\geq 140/90$ on 2 occasions plus one of

- proteinuria
- low platelets
- renal/liver dysfunction
- pulmonary edema
- CNS symptoms



Start IV magnesium sulfate (4g load, then 1g/hr)



Consult OB early and arrange admission or transfer