

THE SAFER MOTHERHOOD

Knowledge Transfer Program

Editor-in-Chief: Professor Sir Sabaratnam Arulkumaran

Pre- eclampsia

Diagnosis and Management



THE GLOBAL LIBRARY OF WOMEN'S MEDICINE

www.glowm.com

WHAT IS PRE-ECLAMPSIA?

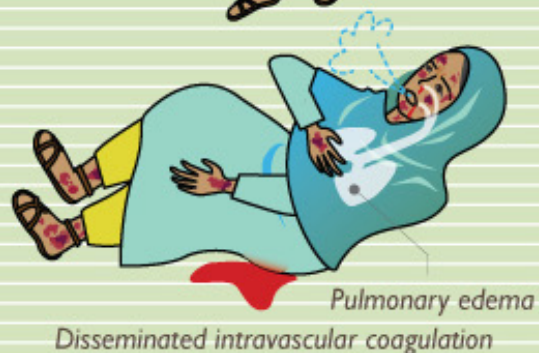
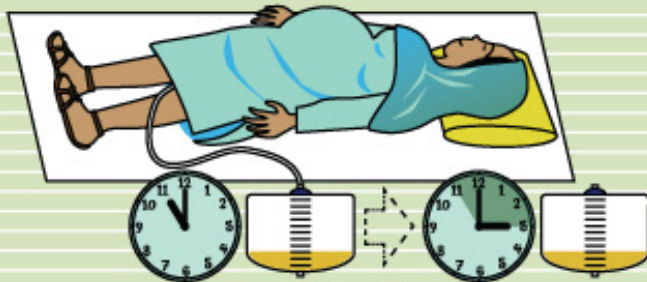
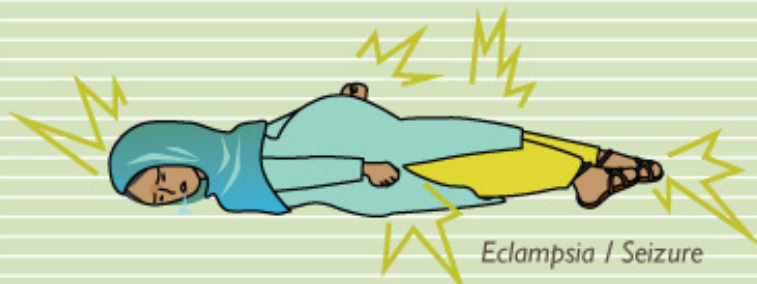
Hypertension in pregnancy :

$BP \geq 140/90$ mmHg

Proteinuria in pregnancy :

$Protein \geq 300$ mg
in 24-hour urine
OR $\geq 1+$ on dipstick

Serious consequences of pre-eclampsia include:



Pre-eclampsia is more than hypertension, proteinuria, and seizures. It is often without any symptoms until the condition deteriorates. The progressive, and unpredictable nature of the disease makes it potentially life-threatening.

Diagnosing

The two primary signs used to diagnose pre-eclampsia are

blood pressure & proteinuria.

Tips

for taking blood pressure measurements accurately:

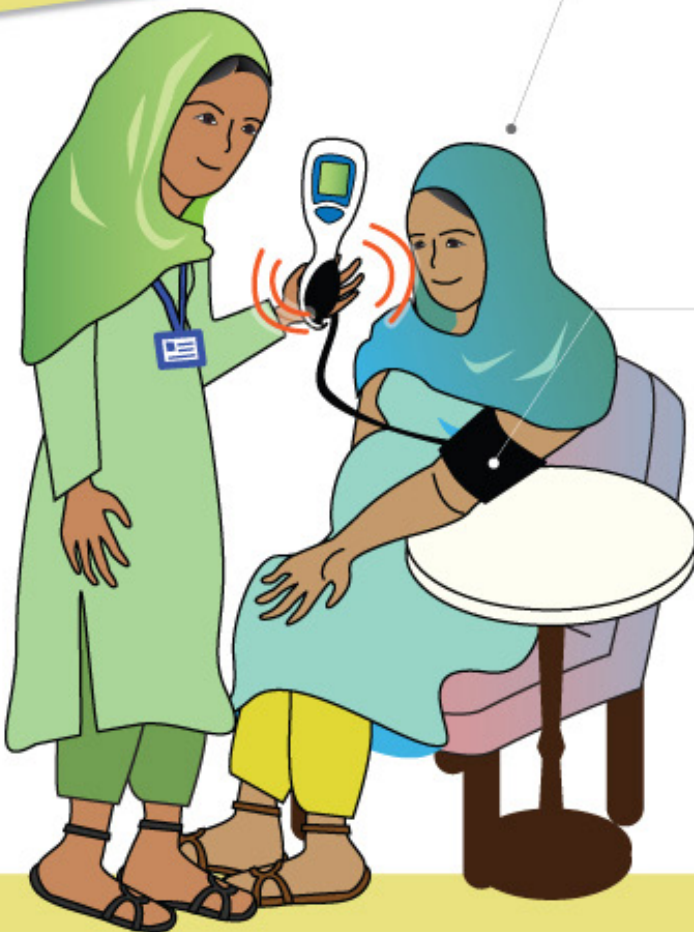
1. Blood Pressure Assessment

Women should be seated **comfortably** with back supported.

The woman should **stay still** for 5 minutes before and during the measurement.

The cuff should be placed around the upper arm, and the arm should be supported at the **level of the woman's heart**.

Measurements should be **repeated at least once**, after a minimum of 1 minute wait, to ensure accuracy.



Community Level Interventions
for Pre-eclampsia

CLIP



2. Proteinuria Assessment



It is best to use a urine sample that is freshly collected directly into the specimen bottle.

Tips

for testing urine sample:

After dipping the protein test strip into the sample, **wait one minute** before reading the result.

Assessing Disease Severity Part I:



Severe hypertension
(very high blood pressure)
is defined as

BP \geq 160/110 mmHg

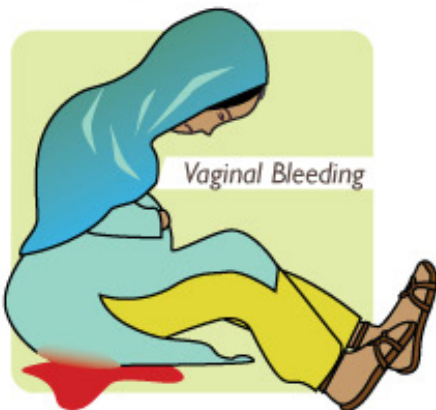
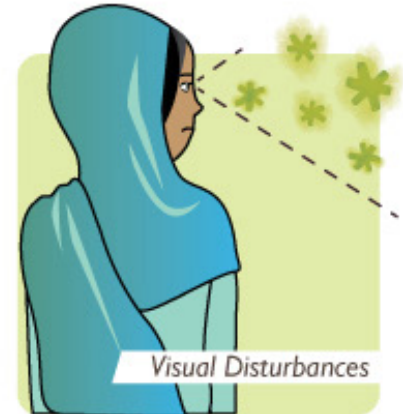
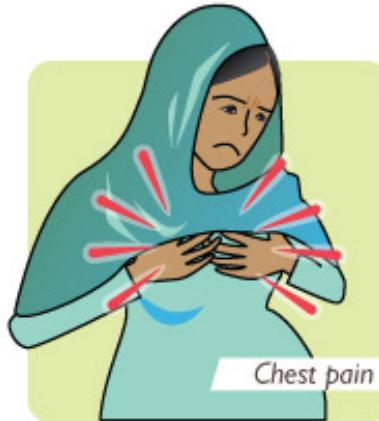


Women with **severe hypertension** are at high risk for **maternal complications**



Women with the highest level of **proteinuria** are at high risk of **stillbirth**

Assessing Disease Severity Part2:



The presence of any of these **symptoms** in a woman with **high blood pressure in pregnancy** indicates that she is **severely ill** and at higher risk of complications such as seizures.

Development of the disorder early in pregnancy is another risk factor for complications of pre-eclampsia.

Therefore, **estimating gestational age early** is an important part of monitoring pregnant women.

Antihypertensive treatment



Acute treatment of severe hypertension should begin **immediately**.

Once blood pressure is **reduced** to the non-severe range (< 160/110 mmHg) ongoing treatment should be initiated using *oral medication*.

Antihypertensive therapy administration instructions by severity of hypertension

severe hypertension

Defined as

BP \geq **160/110** mmHg

Treatment goal: **<160/110mmHg over hours**
(not below 130/80mmHg on antihypertensive therapy)

Oral treatment:

α-Methyldopa

Repeat dose after 3 hr until treatment goal achieved

750 mg

Nifedipine capsules

Repeat dose after 30 minutes until treatment goal achieved

5-10 mg

Nifedipine intermediate-release tablets

Repeat dose after 1 hr until treatment goal achieved

10 mg

Labetalol

Repeat dose after 1 hr until treatment goal achieved

200 mg

Intravenous treatment:

Hydralazine :

Repeat dose after 30 minutes until treatment goal achieved, to a maximum of 20mg

5 mg i.v.

Labetalol :

Repeat dose after 30 minutes until treatment goal achieved, to maximum of 300mg then switch to oral

10-20 mg i.v.

non-severe hypertension

Defined as

BP between **140-159/ 90-109** mmHg

Treatment goal: **<140/90mmHg over days**
(not below 130/80mmHg on antihypertensive therapy)

Oral treatment:

α-Methyldopa

Given 3-4 x daily to a maximum of 2000mg/d

250 mg

Nifedipine intermediate-release tablets (e.g. 'retard' or 'PA')

Given 2 x daily to a maximum of 120mg/d

10-20mg

Labetalol

Given 2-4 x daily to a maximum of 1200mg/d

100-200 mg

Intravenous treatment:

N/A

Eclampsia prevention & treatment

Prevention or Treatment
— of the —
seizures
of eclampsia

NO.1 MgSO₄
is the **best choice** for prevention or treatment

Treatments that are **NOT** recommended:
Phenytoin

Diazepam / lytic cocktail

Recommended treatment or prevention regimens for **MgSO₄** include both

intramuscular *AND* **combined**
intramuscular and intravenous regimens.



MgSO₄	Intramuscular only	Intramuscular and intravenous
Loading dose*	10g i.m. (5g/10mL solution in each buttock)	4g i.v. with 10g i.m. (5g in each buttock)
Maintenance dose*	5g i.m. into alternating buttocks every 4 hours for 24 hours	Either 1-2 g/hr i.v. OR 5g in alternating buttocks every 4 hours for 24 hours

* NOTE: If the women received a loading dose of MgSO₄ in the community (by i.v. / i.m. or i.m. only), maintenance therapy should be initiated if she arrives at the facility within 6hrs. If more than 6hrs has passed since the loading dose was administered in the community, a second loading dose should be administered prior to starting maintenance therapy.

Magnesium Sulphate ($MgSO_4$) intramuscular administration instructions:

1. Explain the reason and procedure briefly to the woman or attendant (as appropriate)

2. Wash hands



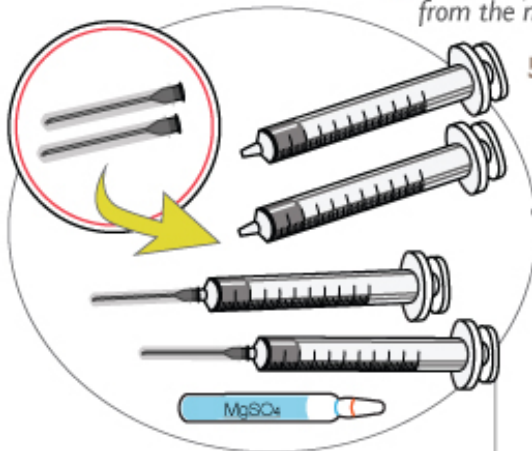
3. Put on gloves



4. Take the pre-prepared syringe from the medicine box

5. Check $MgSO_4$ expiry date and ensure medication is clear and colourless

6. Attach needle to syringe and ensure needle is securely attached (x2)



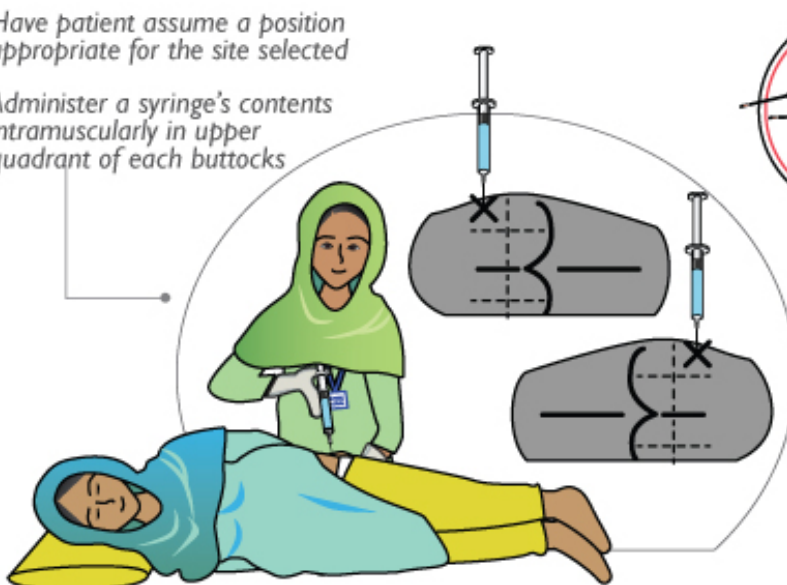
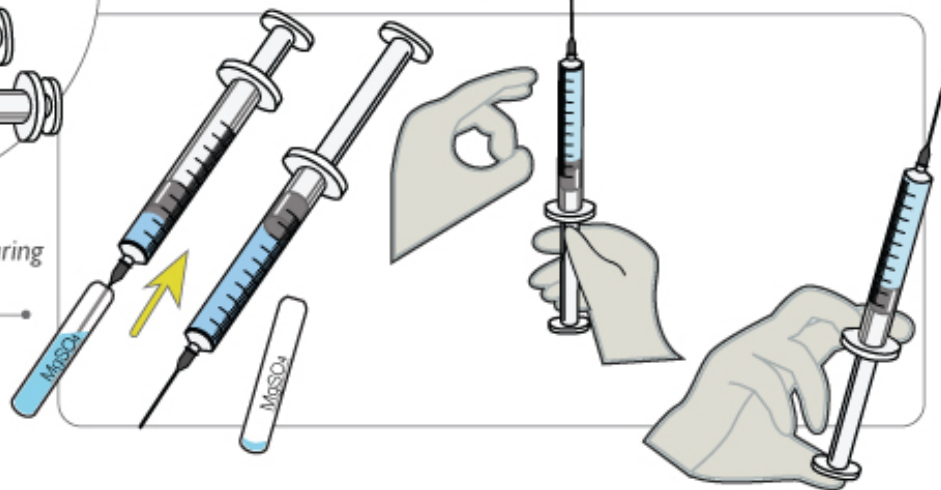
7. Fill each syringe with contents, ensuring 5g $MgSO_4$ / 10mL total volume

8. Clean the injection site with alcohol swab

9. Inspect skin surface for bruises, oedema, or inflammation

10. Have patient assume a position appropriate for the site selected

11. Administer a syringe's contents intramuscularly in upper quadrant of each buttocks



12. Discard needles and syringes in the designated disposal container

13. Apply pressure to the injection sites for 2 minutes

Gestational age at diagnosis

WEEKS	20 wk - viability	viability - 30 wk	30 - 35 wk	35 - 37 wk	≥37 wk
Transfer to referral centre while pregnant	Not necessary. However, centre should be competent with midtrimester termination.	Yes, if stable for transfer	Yes, if stable for transfer. Perinatal outcomes unchanged if transfer occurs postpartum.		
Expectant management	No, as routine. May be attempted close to viability to give fetus a chance.	Yes, as significant perinatal gains without an increase in adverse maternal outcomes. Delivery decision guided by results of maternal and fetal testing. If testing not possible, delivery the safer option.	Yes, due to immediate morbidity and school age issues related to late preterm birth.	No, IOL indicated.	
Corticosteroids for fetal lung maturation	No	Yes	Yes	up to 34+6 wk or 35+6 wk according to local protocol	No
Route of delivery	Vaginal (misoprostol or Foley catheter for labour induction).	Probable Caesarean section, unless intrauterine fetal death.	Vaginal, although fetal, maternal, or uterine status may preclude vaginal delivery.		

Postpartum Patient Care

Severe disease may deteriorate transiently postpartum.

- Maintain surveillance and provide organ system support, as necessary.

Postpartum BP reaches its maximal levels between days 3-6 after delivery.

- If patient is on antihypertensives antenatally, consider maintaining treatment postpartum.
- BP targets can be lower as there are no fetal concerns. In high-income countries, more than 50% of eclampsia occurs postpartum.