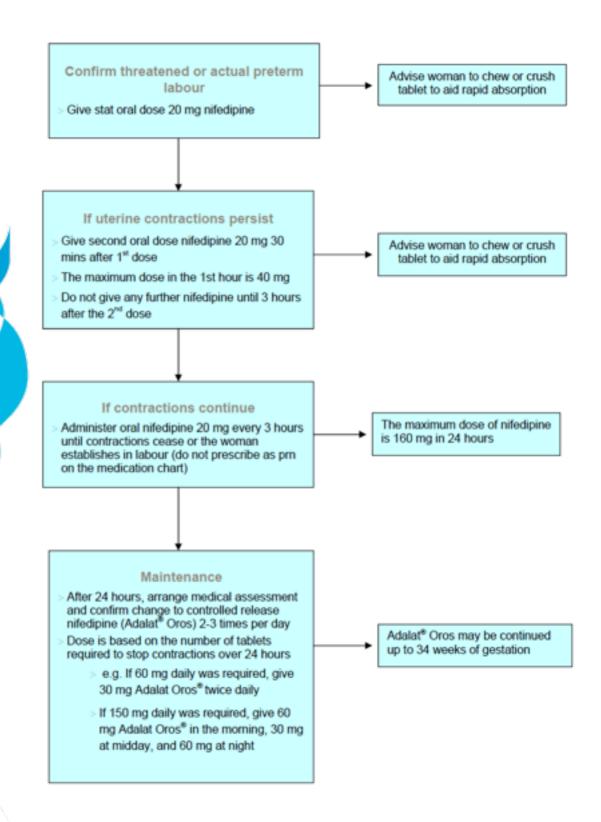
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## Nifedipine for suppression of preterm labour





ISBN number: Endorsed by: Contact: UNKNOWN
SA Maternal & Neonatal Clinical Network
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## **Product information**

- Nifedipine, a dihydropyridine calcium channel blocker, is an effective smooth muscle relaxant with low toxicity. Although nifedipine is known as an antihypertensive drug, the drop in blood pressure in normotensive women after starting tocolytic therapy is significantly more with intravenous salbutamol as compared to nifedipine
  - Adalat®: Absorption of nifedipine is delayed. Maximum plasma concentrations occur in 1.5 4.2 hours. The half life is 6 12 hours
  - Adalat Oros® (controlled release nifedipine) reaches a plateau 6 hours after administration. Adalat Oros® must be swallowed whole
  - Nifedipine is metabolised by the liver and the inactive metabolite is excreted mainly by the kidney
  - In case of urgency ask the woman to chew the tablet and swallow. This does not apply to Adalat Oros<sup>®</sup>. Grapefruit juice inhibits hepatic metabolism
  - Research has shown nifedipine to be a more effective tocolytic agent than betamimetics in prolonging pregnancy for preterm labour (King et al. 2003)
  - Nifedipine is classified as a risk Category C drug by the Australian Drug Evaluation Committee
  - Nifedipine carries the potential for fetal hypoxia associated with maternal hypotension
  - The TGA approved Product information and Consumer Medicine Information says: "Nifedipine is contraindicated throughout pregnancy, and breastfeeding should be stopped first if nifedipine treatment becomes necessary during the breastfeeding period." and "Do not take if you are pregnant or breastfeeding", respectively
  - This guideline should only be used in consultation with specialists who are familiar with the management of preterm labour and the care of preterm infants

### **Indications**

Suppression of

- Threatened Preterm Labour < 34 weeks</p>
- Actual Preterm Labour < 34 weeks</p>

## Contraindications

#### **Maternal**

- > Hypotension (systolic BP less than 90 mmHg) carries the potential for fetal hypoxia
- > Allergy to nifedipine
- Cardiac disease (congestive cardiac failure, aortic stenosis)
- > Hepatic dysfunction

### Fetal

- > Proven intrauterine infection
- > Fetal compromise requiring delivery

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- Placental abruption
- > Severe growth restriction
- > > Lethal fetal anomalies
- Intrauterine fetal death (IUFD)

## Relative contraindications

#### **Maternal**

- Concurrent use of IV salbutamol, transdermal nitrates (GTN)
- Nifedipine and magnesium sulphate (MgSO<sub>4</sub>): Concomitant use of MgSO<sub>4</sub> with nifedipine may result in significant hypotension, and neuromuscular weakness if using the conventional 4 – 6 g IV bolus. An alternative is continuous infusion of MgSO<sub>4</sub> 1 g / hour (King et al. 2003)
- PPROM after adequate steroid cover (48 hours)

#### **Fetal**

- Suspected intrauterine infection
- Preterm labour in the presence of placenta praevia
- Undiagnosed significant vaginal bleeding

## Dosage

Check blood pressure before administering nifedipine

### Confirm threatened or actual preterm labour

Sive stat dose nifedipine 20 mg. The tablet should be chewed or crushed to aid the speed of absorption (grapefruit juice increases the bioavailability of nifedipine by inhibiting its metabolism)

## If uterine contractions persist

- > The second dose of nifedipine 20 mg is given 30 minutes after the first dose. The tablet should be chewed or crushed to maximise speedy absorption
- The maximum dose of nifedipine in the first hour is 40 mg
- > Do not give any further nifedipine until three hours after the second dose

#### If contractions continue

- Administer nifedipine 20 mg every three hours as prescribed, i.e. until the contractions cease or the woman establishes in labour
- Prescribe as written above (do not prescribe as prn)
- > The maximum dose of nifedipine is 160 mg in 24 hours

#### Stop the nifedipine if:

- There is marked hypotension, e.g. systolic < 90 mm Hg</p>
- > Significant dyspnoea

## Observations

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- Maternal baseline BP, TPR, FHR before administering the first dose of nifedipine 20 mg
- > Continue hourly BP and maternal pulse for four hours
- > Check BP before administering nifedipine
- Temperature every 4 hours
- > The rate of observations should be tapered according to the clinical situation
- > Continuous CTG while contracting
- Recommence CTG in the presence of:
  - Regular abdominal pains or tenderness
  - Change in amount, colour of liquor
  - > Antepartum haemorrhage
- > And arrange medical review

#### Side Effects

- > In normotensive women, the effects of nifedipine on BP are minimal
- > Headache
- > Tachycardia, palpitations
- Flushing
- Fatigue
- Dizziness
- Constipation
- Nausea and heartburn.
- > Peripheral oedema secondary to arteriolar vasodilatation
- > Transient rise in liver function test results
- NB: Care with concomitant use of antihypertensive medications (check blood pressure before administering nifedipine)

## Maintenance treatment with nifedipine

- Evidence for the effectiveness of maintenance treatment is lacking (Naik Gaunekar, Crowther 2004)
- After 72 hours, the dosage of nifedipine tablets that was required to stop contractions is given as nifedipine controlled release tablets (Adalat Oros®) 2 – 3 times per day and can be continued to 34 weeks of gestation

#### For example:

- > If 60 mg daily was required, give 30 mg Adalat Oros® twice daily
- If 150 mg daily was required, give 60 mg (Adalat Oros®) in the morning, 30 mg (Adalat Oros®) at midday, and 60 mg (Adalat Oros®) at night

## References

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### Useful reference

RANZCOG College Statement: The use of nifedipine in obstetrics. Available from URL: http://www.ranzcog.edu.au/publications/statements/C-obs15.pdf



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## **Abbreviations**

mm Hg	Millimetres of mercury		
BP	Blood pressure		
IV	Intravenous		
GTN	Glyceryl trinitrate		
et al	And others		
MgSO <sub>4</sub>	Magnesium sulphate		
PPROM	Preterm prelabour rupture of the membranes		
g	Gram/s		
min	minute		
IUFD	Intrauterine fetal death		
TGA	Therapeutic goods administration		
mg	milligram/s		
e.g.	for example		
TPR	Temperature, pulse, respirations		
FHR	Fetal heart rate		
CTG	Cardiotocograph		

## Version control and change history

PDS reference: OCE use only

Version	Date from	Date to	Amendment
1.0	16 June 04	18 Mar 10	Original version
2.0	18 May 10	18 Oct 10	Reviewed
3.0	18 Oct 10	current	



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