WHO recommendation on antenatal corticosteroid therapy in women with preterm prelabour rupture of membranes

17 November 2015

Recommendation

Antenatal corticosteroid therapy is recommended in women with preterm prelabour rupture of membranes and no clinical signs of infection.

(Strong recommendation based on moderate-quality evidence for newborn outcomes and low-quality evidence for maternal outcomes)

Publication history

First published: November 2015

Updated: No update planned

Assessed as up-to-date: November 2015

Remarks

- The use of prophylactic antibiotics should be included as part of standard care for the mother once preterm prelabour rupture of the membranes is confirmed.
- The GDG noted the paucity of evidence on benefits with regard to the duration of membranes rupture due to the lack of such information from trials included in the review. However, the group placed its emphasis on the overall balance favouring benefits over harms of using antenatal corticosteroids in terms of reducing severe adverse neonatal outcomes without evidence of increased risk of infection to the mother or the baby, and with the consideration that a substantial proportion of women at risk of imminent preterm birth would present with ruptured membranes, and therefore made a strong recommendation.
- The GDG cautioned against the use of antenatal corticosteroids for women with prolonged rupture of the membranes and with features of sepsis.
Background

Preterm birth, defined as birth before 37 weeks of gestation, is the single most important determinant of adverse infant outcomes, in terms of survival and quality of life. (1) Globally, it is the leading cause of perinatal and neonatal mortality and morbidity. (2) Preterm infants are particularly vulnerable to complications due to impaired respiration, difficulty in feeding, poor body temperature regulation and high risk of infection. (3-5) With the increasing contribution of neonatal deaths to overall child mortality, it is critical to address the determinants of poor outcomes related to preterm birth to achieve further reductions in child mortality. (6-8)

Infant mortality and morbidity from preterm birth can be reduced through interventions delivered to the mother before or during pregnancy, and to the preterm infant after birth. (9) Interventions can be directed at all women for primary prevention and reduction of the risk of preterm birth (e.g. smoking cessation programme) or aimed at minimizing the risk in women with known risk factors (e.g. progestational agents, cervical cerclage). (10) However, the most beneficial set of maternal interventions are those that are aimed at improving outcomes for preterm infants when preterm birth is inevitable (e.g. antenatal corticosteroids, magnesium sulfate and antibiotic prophylaxis). (9) Special care of the preterm newborn to prevent and treat complications of prematurity is also critical to newborn survival. In high-income countries, reductions in mortality rates in infants that were born preterm have been driven largely by improved care and, more importantly, by appropriate policy changes.

Methods

The recommendations were developed using standard operating procedures in accordance with the process described in the WHO handbook for guideline development (11). Briefly, these included (i) identification of priority questions and critical outcomes, (ii) retrieval of the evidence, (iii) assessment and synthesis of evidence, (iv) formulation of recommendations, and (v) planning for the dissemination, implementation, impact evaluation and updating of the guideline.

The scientific evidence underpinning the recommendations was synthesized using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach (12). Up-to-date systematic reviews were used to prepare evidence profiles for the priority questions. WHO then convened a Technical Consultation in May 2014 where an international group of experts – the Guideline Development Group (GDG) – formulated and approved the recommendations based on the evidence profiles.

In November 2014, an online consultation of the GDG was conducted to review and revise the recommendations in the light of the findings of a large implementation trial of antenatal corticosteroids in low-resource countries.

Further information on procedures for developing this recommendation are available here.

Recommendation question

For this recommendation, we aimed to answer the following questions:

- Among pregnant women at risk of imminent preterm birth (P), is antenatal corticosteroid therapy (I), compared with no antenatal corticosteroid therapy (C), effective in reducing adverse newborn outcomes (O)?
• Which population of pregnant women should be offered antenatal corticosteroids? (considering the gestational age at presentation or birth; interval between presentation and anticipated birth; single and multiple birth; status of amniotic membranes; and women undergoing elective caesarean section in late preterm)

• Which population of pregnant women should not be offered antenatal corticosteroids? (considering conditions where there are concerns that associated risks may outweigh benefits: women with diabetes mellitus, hypertensive disorders, chorioamnionitis and growth-restricted babies)

• Which corticosteroids (and regimens) should be used for eligible women?

• Should repeat course(s) of corticosteroids be offered to a woman who has completed a course of corticosteroid but remains at risk of preterm birth 7 days or more after the initial treatment?

Evidence summary

**Antenatal corticosteroids versus placebo or no treatment (preterm prelabour rupture of membranes)**

In a Cochrane review (13), the effects of antenatal corticosteroids were examined in a subgroup of women with PPROM.

**Maternal outcomes**

*Severe morbidity or death:* No significant differences were observed between groups for maternal death, chorioamnionitis or puerperal sepsis in mothers when the first dose of corticosteroids was given to women with PPROM or prolonged rupture of membranes (> 24 hours).

**Infant outcomes**

*Fetal and neonatal death:* Combined fetal and neonatal deaths were significantly reduced among infants exposed to antenatal corticosteroid and born following PPROM at the time of first dose (RR 0.62, 95% CI 0.46–0.82; 4 studies, 733 infants), but not following prolonged rupture of membranes > 24 hours (RR 0.77, 95% CI 0.51–1.17; 2 studies, 508 infants) or > 48 hours (RR 0.93, 95% CI 0.57–1.51; 1 study, 255 women). As with previous outcomes, this reduction was due to the contribution of reduced neonatal mortality among corticosteroid-exposed infants (RR 0.61, 95% CI 0.46–0.83; 8 studies, 1024 infants), while no reduction in fetal deaths was observed for any of these subgroups.

*Severe neonatal morbidity:* RDS was significantly reduced in infants whose mothers received corticosteroids at the time of PPROM (RR 0.68, 95% CI 0.57–0.83; 12 studies, 1129 infants), as was cerebroventricular haemorrhage (RR 0.47, 95% CI 0.28–0.79; 5 studies, 895 infants), necrotizing enterocolitis (NEC) (RR 0.39, 95% CI 0.18–0.86; 4 studies, 583 infants), and duration of mechanical ventilation (MD -3.50 days, 95% CI -5.12 to -1.88 days; 1 study, 165 infants). No significant differences were observed for neonatal infection, systemic infection in the first 48 hours, or need for mechanical ventilation or continuous positive airway pressure (CPAP).

Further information and considerations related to this recommendation can be found in the WHO guidelines, available at:

http://apps.who.int/iris/bitstream/handle/10665/183037/9789241508988_eng.pdf?sequence=1
Implementation considerations

- The successful introduction of this recommendation into national programmes and health-care services depends on well-planned and participatory consensus-driven processes of adaptation and implementation. The adaptation and implementation processes may include the development or revision of existing national guidelines or protocols based on this recommendation.
- The recommendation should be adapted into a locally appropriate document that can meet the specific needs of each country and health service. Any changes should be made in an explicit and transparent manner.
- A set of interventions should be established to ensure that an enabling environment is created for the use of the recommendations, and that the behaviour of the healthcare practitioner changes towards the use of this evidence-based practice.
- In this process, the role of local professional societies is important and an all-inclusive and participatory process should be encouraged.

Research implications

The GDG identified these priority questions related to antenatal corticosteroids for imminent preterm birth recommendations:

- What are the long-term outcomes of all infants exposed to antenatal corticosteroids (including term infants)?
- What strategies can effectively and safely increase the use of corticosteroids in low- and middle-income country (LMIC) settings to improve outcomes?
- What are the effects of antenatal corticosteroid at different gestational ages at birth (using independent patient data analysis)?
- Assessment of coverage of antenatal corticosteroids before and after guideline implementation (and associated reduction in neonatal mortality).
- Assessment of implementation strategies and monitoring of adverse events (in LMIC settings).
- What are the effects of task shifting in the context of antenatal corticosteroid administration (e.g. using the first dose in the community followed by referral to a health-care facility)?
- Are there differences in the pharmacokinetic properties of betamethasone acetate versus betamethasone phosphate (consider using available data in settings where they are routinely used)?
- What is the impact of antenatal corticosteroid administration among mothers with evidence of infection who also receive appropriate antibiotic therapy on both maternal and neonatal outcomes?
- What is the minimum effective dose of corticosteroids to achieve fetal lung maturation and other improved outcomes?
- What is the minimum effective dose required for repeat courses of antenatal corticosteroids?
- What is the most effective regimen and dose for antenatal corticosteroids?
- In what contexts can antenatal corticosteroids be used safely and effectively in low-income countries?

Related links

WHO recommendations on interventions to improve preterm birth outcomes (2015) –[full document](#) and [evidence tables](#)
Managing Complications in Pregnancy and Childbirth: A guide for midwives and doctors

Pregnancy, Childbirth, Postpartum and Newborn Care: A guide for essential practice

WHO Programmes: Sexual and Reproductive health

Maternal Health

Infant, Newborn Health

Supporting systematic review:


References
