



Preterm Premature Rupture of Membranes (PPROM)

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Course Description:

Leaking fluid? We have problems! Preterm Premature Rupture of Membranes (PPROM) continues to be a leading cause of neonatal morbidity and mortality. This module will help to develop a knowledge base to further the care of a woman with PPROM.

Approximate Time to Complete: 45 minutes



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In this course the participant will:

- Recognize the risk factors for PPROM.
- The signs and symptoms of PPROM will be recognizable after this module.
- Better understand the physical exam of a woman suspected to have PPROM.
- Have a better understanding of the serious infections that can occur with PPROM patients.
- Understand possible complications for a woman and her developing fetus when she is faced with PPROM.
- Formulate the plan of testing a mother presenting with signs and symptoms of PPROM to help determine if the membranes have ruptured prematurely.
- Be able to explain the controversy with treatment tracks and understand the nuances to the treatment options.
- Gain knowledge on the medication regimens for PPROM based on the gestational age of the fetus.



-  Background Information
 -  Definition
 -  Risk Factors
-  Diagnosis
 -  Diagnosing PPROM
 -  Diagnosing PPROM - Clinical Course
 -  Complications with PPROM
 -  Risks of PPROM
 -  Diagnosing PPROM
 -  Differential Diagnosis
-  Management
 -  Management
 -  Initial Approach
 -  Assessment
 -  Management Decisions
 -  Intervention
 -  Management
 -  Potential Fetal Consequences
 -  Special Circumstances
-  Delivery
 -  Delivery
 -  Cervical Ripening
-  Summary
 - 



Premature rupture of membranes (PROM)

Defined as membrane rupture before the onset of uterine contractions (also called pre-labor rupture of membranes)

Preterm PROM (PPROM)

PROM before 37 weeks 0 days of gestation.

Occurrence of PPROM

PPROM occurs in three percent of pregnancies, but is responsible for or associated with about one-third of preterm births [1].



PPROM Risk Factors

- These risk factors are similar to those for preterm labor (PTL) (Table 1), but most patients have no identifiable risk factors.
- A history of PPROM in a previous pregnancy, genital tract infection, antepartum bleeding, and cigarette smoking have a particularly strong association with PPROM [2].



Click on the table to view a larger version.

Bacteruria	No partner
Periodontal disease	Low socioeconomic status
Placenta previa	Anxiety
Placental abruption	Depression
Vaginal bleeding, especially in more than one trimester	Life events, divorce, separation, death
Previous PTD	Abdominal surgery during pregnancy
Substance abuse	Occupational issues
Smoking	upright postures
Maternal age (younger than 18 or older than 40)	use of industrial machines
African-American race	physical exertion
Poor Nutrition and low body mass index	mental or environmental stress related work
Inadequate prenatal care	working conditions
Anemia (hemoglobin <10g/dL)	Multiple gestation
Excessive uterine contractions	Polyhydramnios
Low level of educational achievement	Uterine anomaly
Genotype	DES induced uterine changes
Fetal anomaly	History of second trimester abortion
Fetal growth restriction	History of cervical surgery
Environmental factors (i.e. heat, air pollution)	Premature cervical dilatation or effacement (shortened cervix)
STI's	

PPROM Risk Factors



Previous PPROM

- PPROM when evaluated in studies has supported a strong risk for recurrence.
- For example, the Preterm Prediction Study, a large prospective study conducted by the National Institute of Child Health and Human Development (NICHD) Maternal-Fetal Medicine Units (MFMU) Network, observed that women with a history of PPROM had a 13.5 percent rate of PPROM in a subsequent pregnancy compared to 4.1 percent in women with no such history [3].
- Others have reported recurrence rates as high as 32 percent [4].
- Women with a history of PPROM are at risk for recurrent PPROM or preterm birth (PTB) without PPROM [5,6].



Slide 2 of 4



PPROM Risk Factors

Genital Tract Infection

- Genital tract infection is the single most common identifiable risk factor for PPRM.
- Three lines of epidemiologic evidence supports this association:
- Women with PPRM are significantly more likely than women with intact membranes to have pathogenic microorganisms in the amniotic fluid.
- Women with PPRM have a significantly higher rate of histologic chorioamnionitis than those who deliver preterm without PPRM.
- Lower genital tract infections, particularly bacterial vaginosis, has been associated with higher rates of PPRM than uninfected women [7].
- There is a strong association between bacterial colonization of the lower genital tract and PPRM.
- Lower genital tract microorganisms can colonize and produce phospholipases. The phospholipases stimulate prostaglandin production and can lead to uterine contractions.
- The host's immune response to endocervical and fetal membrane bacterial invasion can lead to production of inflammatory mediators which will cause weakening of the fetal membranes and can cause in PPRM [7].
- The host's immune and inflammatory response can be genetically regulated and play a role is susceptibility and response to infections that are associated with PPRM.



Slide 3 of 4



PPROM Risk Factors



- **Antepartum bleeding**
- When antepartum bleeding occurs in the first trimester, there is a statistically significant increase in the risk of PPRM [8].
- Bleeding in more than one trimester increases the risk of PPRM by 3-to-7 fold [2,9,10].
- **Cigarette smoking**
- Among smokers, the risk of PPRM climbs by 2-to-4 fold compared to non-smokers [10].
- This risk continues after other treatment for PPRM has been provided, including infection.



Slide 4 of 4



PPROM is noted to have a classic presentation of a sudden gush of clear or pale yellow fluid from the vagina.

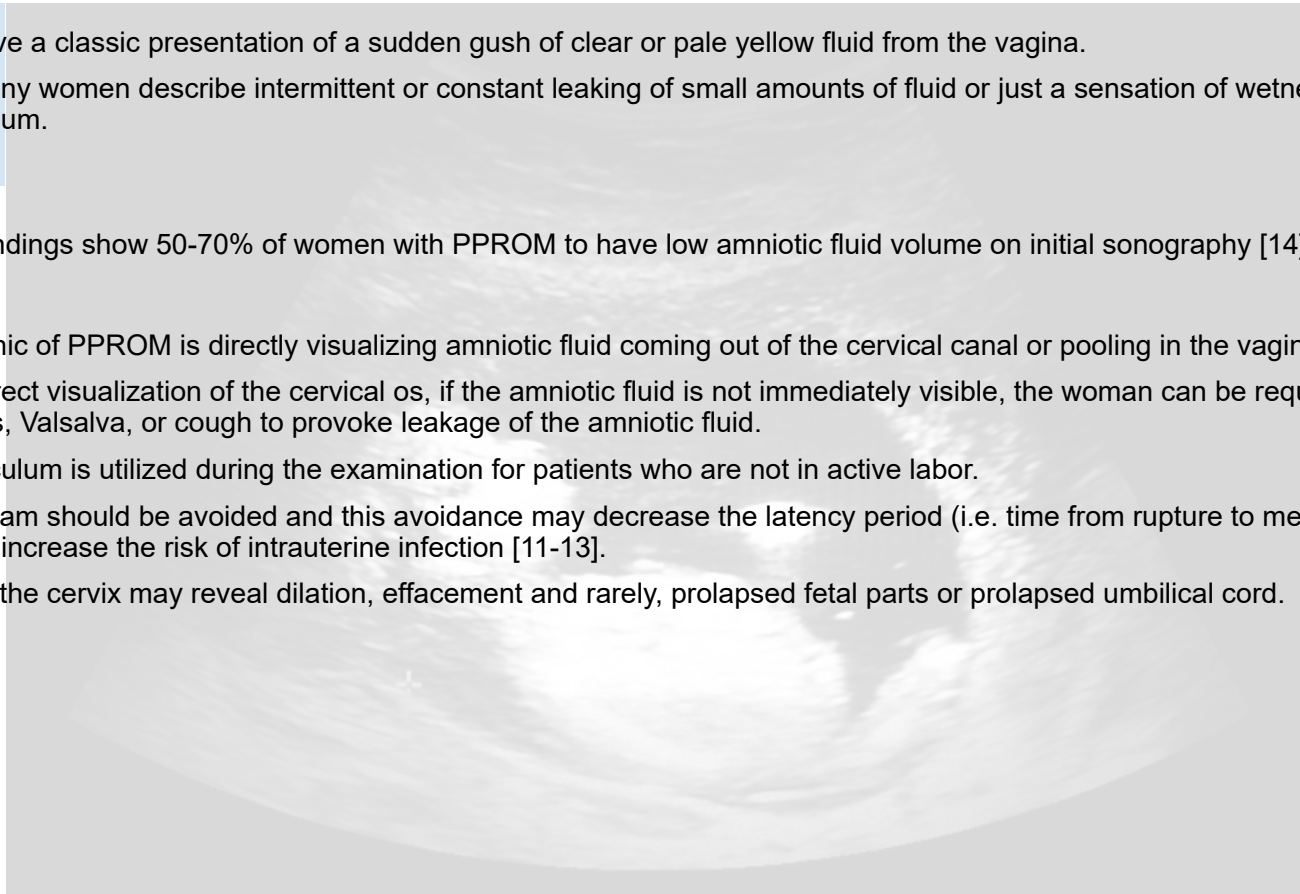
The problem is that many women describe intermittent or constant leaking of small amounts of fluid or just a sensation of wetness within the vagina or on the perineum.

On ultrasonography:

- Ultrasound findings show 50-70% of women with PPRM to have low amniotic fluid volume on initial sonography [14].

Physical Exam:

- Pathognomonic of PPRM is directly visualizing amniotic fluid coming out of the cervical canal or pooling in the vaginal fornix.
- During the direct visualization of the cervical os, if the amniotic fluid is not immediately visible, the woman can be requested to push on her fundus, Valsalva, or cough to provoke leakage of the amniotic fluid.
- A sterile speculum is utilized during the examination for patients who are not in active labor.
- The digital exam should be avoided and this avoidance may decrease the latency period (i.e. time from rupture to membranes to delivery) and increase the risk of intrauterine infection [11-13].
- Inspection of the cervix may reveal dilation, effacement and rarely, prolapsed fetal parts or prolapsed umbilical cord.





1

2

3



*Click each numbered tab
to see more information.*



1

- The majority of pregnancies with PPROM deliver within one week of membrane rupture.

2

- A randomized trial of PPROM at 24-32 weeks showed the median time to delivery of 239 group B streptococcal (GBS) negative women managed expectantly with prophylactic antibiotics was 6.1 days along with the following observations:
 - 27 percent delivered within 48 hours
 - 56 percent delivered within 7 days
 - 76 percent delivered within 14 days
 - 86 percent delivered within 21 days [15]

3



1

2

3

- Of interest, the duration of latency inversely correlates with gestational age at membrane rupture [16].
- Sealing of the membranes is associated with a more favorable outcome, however, cessation of fluid leakage is rare, except in women with PPROM related to amniocentesis [17].
- Unfortunately, the fetus and neonate are at greater risk of PPROM-related morbidity and mortality than the mother ([Table 2](#)).



1

2

3

- The morbidity associated with prematurity varies with gestational age and is higher in the setting of chorioamnionitis [18].
- There is an increased risk of neurodevelopmental impairment when the fetus is exposed to intrauterine inflammation [19].
- About one-third of PPROM patients develop potentially serious infections, such as intraamniotic infection (chorioamnionitis and funisitis), endometritis, or septicemia.
- Endometritis occurs more often following a cesarean than a vaginal delivery.
- Infection occurs at a higher rate the earlier the gestational age [19, 20].



Click on each potential complication as it appears to learn about the potential Offspring and Maternal consequences.

Pregnancy Complication	Potential Consequences for Offspring	Potential Maternal Consequences
Intrauterine infection		





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Intrauterine infection	Neonatal Sepsis Long-term neurodevelopmental abnormalities, particularly cerebral palsy	Postpartum endometritis
Umbilical cord compression		





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Umbilical cord compression	Fetal asphyxia	
Oligohydramnios		





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Oligohydramnios	Limb restriction deformities and pulmonary hypoplasia (primarily with severe oligohydramnios in the early to mid second trimester), occurring rarely when membranes rupture after 23 weeks.	
Fetal malpresentation		





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Fetal malpresentation		Cesarean delivery
Umbilical cord prolapse		





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Fetal malpresentation		Cesarean delivery
Umbilical cord prolapse	Fetal asphyxia	Cesarean delivery
Placental abruption		





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Umbilical cord prolapse	Fetal asphyxia	Cesarean delivery
Placental abruption	Fetal asphyxia	Cesarean delivery Coagulopathy
Preterm Birth		





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Umbilical cord prolapse	Fetal asphyxia	Cesarean delivery
Placental abruption	Fetal asphyxia	Cesarean delivery Coagulopathy
Preterm Birth	Morbidity of prematurity	

Increased risk of placental abruption and prolapsed umbilical cord occurs with PPROM.

- Placental abruption occurs in 2 to 5 percent of pregnancies complicated by PPROM [21-24].
- Placental abruption risk increases even further with an increase of 7-9 fold in PPROM pregnancies with intrauterine infection or oligohydramnios present [22,23].
- Placenta abruption may be the event that either causes PPROM or the consequence of PPROM.

It is common to have fetal malpresentation and reduced amniotic fluid volume at a preterm gestational age.

- The cord prolapse risk is especially high, up to 11% in one study [25], when both non-vertex fetal presentation occurs with PPROM.
- This malpresentation may also increase the risk of abruption, infection, and fetal death in-utero [26].

When PPROM occurs early, is severe and with prolonged oligohydramnios, it can be associated with pulmonary hypoplasia, facial deformation, and orthopedic abnormalities.

- These complications are much more likely when membrane rupture occurs under 23 weeks of gestation.



Nitrazine

Ferning

Ultrasonography

**Instillation of
Indigo Carmine**

AmniSure vs Actim PROM

AmniSure

Actim PROM

fFN

- Generally, the diagnosis of PPROM occurs clinically and is based on the visualization of amniotic fluid in the vagina of a woman who presents with a history of leaking fluid.
- When the diagnosis is uncertain, laboratory tests are utilized.



Click each term to the left to learn more about diagnosing PPROM.

Nitrazine**Ferning****Ultrasonography****Instillation of
Indigo Carmine****AmniSure vs Actim PROM****AmniSure****Actim PROM****fFN**

- After visualization, if PPRM is not obvious, the diagnosis can be confirmed by testing the pH of the vaginal fluid utilizing nitrazine paper or like pH indicator product.
- The typical pH range of amniotic fluid is 7.0 to 7.3 and much higher compared to normal acidic vaginal pH of 3.8 to 4.2 [27].
- In up to five percent of testing, a false negative or false positive result occurs.
- The false negative results can occur when leaking is intermittent or the amniotic fluid is diluted by other vaginal secretions.
- On the other hand, false positive results may occur in the presence of alkaline vaginal fluids, such as blood, seminal fluid, or soap.
- When the urine has infection of proteus species, the pH of the urine may be elevated near 8.0.

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- Another confirmatory test is the presence of arborization, also called ferning.
- To view the arborization, fluid is obtained from the posterior vagina fornix in a woman suspected of having PPROM, swabbing this onto a glass and allowing it to dry for at least ten minutes.
- The dried amniotic fluid produces a delicate ferning pattern, in contrast to the thick and wide arborization pattern of dried cervical mucus.
- False positive fern testing may occur by well estrogenized cervical mucus or a fingerprint on the microscope slide.
- Insufficient amniotic fluid transferred from the swab to the slide or heavy contamination with vaginal discharge or blood can lead to false negative results.



Nitrazine

Ferning

Ultrasonography

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fFN

- In equivocal cases, ultrasound can be performed to look for a reduction in amniotic fluid volume.



Nitrazine

Ferning

Ultrasonography

**Instillation of
Indigo Carmine**

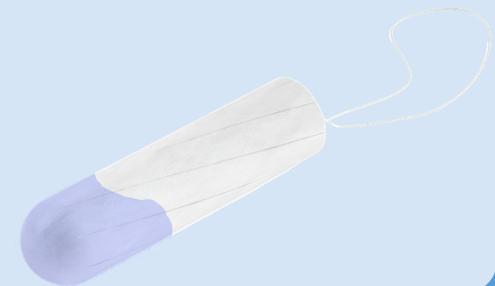
AmniSure vs Actim PROM

AmniSure

Actim PROM

fFN

- When there is continued uncertainty of PPROM, instillation of indigo carmine into the amniotic cavity may lead to a definitive diagnosis.
- Utilizing ultrasound guidance, 1 mL of indigo carmine in 9 mL of sterile saline is injected transabdominally into the amniotic fluid.
- A tampon is placed in the vagina.
- When the tampon is removed it is examined for blue staining, typically about 20 minutes later, and if present PPROM is confirmed.
- Of note, the maternal urine will also turn blue so this finding should not be confused with leaking of amniotic fluid.
- Alternatives to indigo carmine include sodium fluorescein and phenolsulfonphthalein where available.



Nitrazine

Ferning

Ultrasonography

**Instillation of
Indigo Carmine**

AmniSure vs Actim PROM

AmniSure

Actim PROM

fFN

- There are commercially available tests for diagnosing ROM, two of which are:
- Amnisure which is placental alpha microglobulin-1 protein assay (PAMG-1)
- Actim PROM, an insulin growth factor binding protein 1 (IGFBP-1)
- A meta-analysis from 2013 of prospective observational or cohort studies investigating AmniSure was more accurate than Actim PROM for diagnosing ROM in all patient populations (i.e. known or unknown rupture status) [30].
- A subsequent randomized trial reported similar findings [31].

Nitrazine

Ferning

Ultrasonography

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AmniSure vs Actim PROM

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- AmniSure (placental alpha microglobulin-1 protein assay) is rapid slide test using immunochromatography methods to detect trace amounts of placental alpha microglobulin-1 protein in vaginal fluid.
- A significant advantage to AmniSure testing is the testing is not affected by semen or trace amounts of blood.
- There is a commercially available kit that is obtained by the provider at the point of care.
- The kit has a sterile swab, this is inserted into the vagina for one minute, then placed into a vial containing a solvent for one minute and then an AmniSure test strip is dipped into the vial.
- The result is revealed by the presence of one or two lines within the next 5-10 minutes:
- No visible line is an invalid result
- One visible line is a negative result
- Two visible lines is a positive result
- The sensitivity ranged from 94.4-98.9 percent with a specificity range of 87.5-100 percent in large studies [32-36].
- There were three false positives in one study and the authors hypothesized these have occurred due to a small leak that sealed over [33].
- The cost of the test is relatively high, often limiting the use to cases where the diagnosis remains uncertain after physical exam with ferning and nitrazine testing.

Nitrazine**Ferning****Ultrasonography****Instillation of
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- In problematic cases, the identification of insulin-like growth factor 1 (IGFBP-1) may be of value in confirming the diagnosis of PPROM.
- This protein, insulin-like growth factor binding protein 1, is secreted by decidual and placental cells. It has a very high concentration in amniotic fluid compared to other body fluids.
- The method is easy to use with a immunochromotography dipstick method (Actim PROM) is available in some countries for use at the bedside to detect IGFBP-1 in vaginal secretions.
- This test is popular in Europe, but is not widely used in the United States.
- Two blue lines on the dipstick indicate a positive test.
- The test is not affected by the presence of infected vaginal secretions, urine, semen, or small amounts of blood.
- This test is most accurate when performed as soon as possible after PROM.
- Sensitivity ranges from 95-100 percent in detecting ruptured membranes with specificity ranging from 93-98 percent, and positive predictive values approach 98 percent [35, 37-40].
- This test is very helpful in identifying women likely to deliver within seven days.

Nitrazine

Ferning

Ultrasonography

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fFN



A negative fetal fibronectin result strongly supports the absence of membrane rupture.

A positive result only indicates disruption of the interface between chorion and decidua, which can occur with intact membranes. [41]

- There are other causes of vaginal wetness
 - Urinary incontinence
 - Vaginal discharge
 - Perspiration
 - When the clinical and laboratory findings for PPROM are negative, these other causes should be considered.
 - A mild reduction in amniotic fluid volume, by ultrasound findings, is non-specific and related to many etiologies including PPROM.
- Highly suggestive of PPROM is the finding on ultrasound of anhydramnios or severe oligohydramnios, combined with a characteristic history.
 - Although, renal agenesis, obstructive uropathy or severe utero-placental insufficiency may be the etiology to marked reductions in amniotic fluid volume.



- Some of the most controversial issues in perinatal medicine come from contentions in treating PPRM.
- Points of contention include:
 - Expectant management versus intervention
 - Use of tocolytics
 - Duration of administration of antibiotic prophylaxis
 - Timing of administration of antenatal corticosteroids
 - Methods of testing for maternal/fetal infection
 - Timing of delivery
 - Treatment of women from 23-37 weeks gestation who have PPRM will be reviewed.
 - Treatment of previable PPRM and ROM at term are topics beyond the scope of this program.





PPROM management in women is based upon consideration of several factors, which are assessed at presentation:

- Gestational age
- Presence or absence of maternal and/or fetal infection
- Presence or absence of labor
- Fetal presentation
- Fetal well-being
- Fetal lung maturity
- Cervical status (by visual inspection)
- Sterile speculum (by visualization with a sterile speculum)
- Availability of neonatal intensive care

Assessment



- Nitrazine and fern
- Placental alpha microglobulin-1 protein assay (AmniSure)
- Once confirmed PPROM then consider:
 - Complete blood count
 - Culture for *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, *Trichomonas vaginalis*, and bacterial vaginosis.
- Fetal lung maturity testing
 - Lamellar body count in amniotic fluid
 - Lecithin:sphingomyelin (L/S) ratio
- Rectovaginal culture for group B streptococcus
- Ultrasound to determine:
 - Fetal growth
 - Fetal position
 - Residual amniotic fluid volume
 - Fetal anatomy
- Biophysical profile
- Cardiotocography monitoring for fetal heart rate (non-stress test) and uterine contraction frequency

- A key factor in managing PPRM is whether to induce labor (or perform a cesarean) or to manage expectantly.
- An immature fetus may benefit by prolonging the pregnancy which could result in significant reduction in gestational aged related morbidity. However, this benefit needs to balance with the risks of PPRM associated complications and their sequelae ([Table 2](#)).
- When intratuterie infection, abruptio placentae, nonreassuring fetal testing, or high risk of cord prolapse is present or suspected, then expeditious delivery of these women is appropriate.
- With each of these situations, fetal well being can quickly deteriorate during expectant management and there are no therapeutic interventions available other than delivery.
- When these complications are absent, interventions leading to delivery are not indicated, until 34 weeks, then proceeding with delivery is appropriate.
- An algorithm to manage women with PPRM at 26-37 weeks is shown ([Algorithm 1](#)).
- Several aspects of management will be discussed, however, a detailed analysis of the nuances in managing women with PPRM is beyond the scope of this program.

- The optimal time to intervene varies by facility and depends on the balance between morbidity related to prematurity and morbidity related to complications from PPROM.
- American College of Obstetricians and Gynecologists (ACOG) suggests delivery of all patients at 34 weeks 0 days gestation [42].
- Between 28-37 weeks gestation, meta analysis of randomized trials and subsequent randomized trials, have not provided conclusive evidence favoring induction or expectant management of women with PPROM [43-46].
- The data is limited for analysis due to heterogeneity among the trials.
- Example are fetal lung maturity not being consistently determine and thus not a factor in selecting patients who may or may not benefit from expectant management.
- Another example that could not be assessed is the administration of prophylactic antibiotics because there was not a standard practice and patient level data were not analyzed.
- The last concern involves the power of the trials; the trials have been underpowered to detect meaningful measures of neonatal and maternal morbidity.

Administration of Antenatal Corticosteroids

Antibiotic Therapy

Prophylaxis

Drug Regimen

Chemoprophylaxis for GBS

Tocolysis

Supplemental Progesterone

Hospitalization versus Home Care

Maternal Monitoring

Fetal Monitoring

- When women have a gestation between 23-34 wks, have been diagnosed with PPROM, then a course of corticosteroids should be given.
- Two systematic reviews of randomized trials [47,48] provide data supporting antenatal glucocorticoid treatment showing reduced neonatal death, respiratory distress syndrome (RDS), intraventricular hemorrhage (IVH), necrotizing enterocolitis (NEC), and duration of neonatal respiratory support without an increase in either maternal nor neonatal infection rates.
- Mean risk reduction for these adverse events ranged from 30 to 60 percent.
- The use of rescue therapy is controversial as available data on potential benefits and harms are not definitive in this setting [49-51]
- It is reasonable to provide a single dose of "rescue" therapy if the woman is clinically estimated to be at high risk of delivery within the next seven days, at least two weeks have passed since the initial course of antenatal corticosteroids, and the initial course was given under 28 weeks gestation.
- The research is inconsistent regarding the effect of PPROM on fetal pulmonary maturation.
- The discordance may be due to failure to adjust for factors affecting neonatal respiratory function such as [52]:
 - Mode of delivery
 - Presence or absence of labor
 - Gestational age
 - Duration of latency



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 - Mode of delivery
 - Presence or absence of labor
 - Gestational age
 - Duration of latency
 - Comorbidities



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- The indication for antibiotics are to prolong latency and to reduce the risk of early onset neonatal group B streptococcal (GBS) infection, when present.
- Prophylactic antibiotics are given for seven days when the woman is less than 34 weeks of gestation at the time of membrane rupture.
- Targets of antibiotic therapy include:
 - Ampicillin and Amoxicillin
 - group B streptococcus
 - many aerobic gram-negative bacilli
 - some anaerobes
 - Azithromycin
 - Ureaplasma for chorioamnionitis with PPROM
 - Chlamydia trachomatis (cause of neonatal conjunctivitis and pneumonitis)



**Administration of Antenatal
Corticosteroids****Antibiotic Therapy****Prophylaxis****Drug Regimen****Chemoprophylaxis for GBS****Tocolysis****Supplemental Progesterone****Hospitalization versus Home Care****Maternal Monitoring****Fetal Monitoring**

- Infection may be both a cause and consequence of PPROM, thus making antibiotics a key component to her treatment regimen.
- Spontaneous PTL may be caused by the infection and may be the indication to proceed with medically - indicated preterm delivery.
- Goals of antibiotic therapy include: reduction in frequency of maternal and fetal infection thereby delaying the onset of preterm labor, prolonging latency, and the need for preterm birth.
- Studies underscore the importance of reducing infection by suggesting a relationship between chorioamnionitis, duration of membraned rupture and development of cerebral palsy or neurodevelopment delay.
- Compared with placebo or no treatment, antibiotic use was associated with significant reductions in:
 - Chorioamnionitis
 - Babies born within 48 hours
 - Neonatal infection
 - Use of surfactant
 - Neonatal oxygen therapy



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- group B streptococcus
- many aerobic gram negative bacilli
- some anaerobes
- Azithromycin for:
 - mycoplasma (an important cause of chorioamnionitis with PPROM)
 - Chlamydia trachomatis (cause of neonatal conjunctivitis and pneumonia)
 - abnormal cerebral ultrasound prior to hospital discharge
- The regimen discussed is similar to the National Institute of Child Health and Human Development (NICHD) Maternal-Fetal Medicine Units (MFMU) Network trial.
 - This trial studied antibiotic therapy for reduction of infant morbidity after PPROM with IV ampicillin 2 g every 6 hours and erythromycin 250 mg every 6 hours for 48 hours FOLLOWED by oral amoxicillin 250 mg every 8 hours and erythromycin 333 mg every 8 hours for five days [16]. This is recommended by ACOG.
 - Due to the ease of administration, improved gastrointestinal tolerance, favorable cost profile and similar efficacy azithromycin is recommended in lieu of a multiple-day course of erythromycin.
 - In a retrospective study the two regimens ampicillin + erythromycin versus ampicillin + azithromycin, were reviewed in women with PPROM and resulted in similar pregnancy and neonatal outcomes [54]:
 - Length of latency
 - Mean birth weight
 - Rates of chorioamnionitis
 - Cesarean delivery
 - Low Apgar score
 - Neonatal sepsis
 - Neonatal respiratory distress syndrome (RDS)



Administration of Antenatal Corticosteroids

Antibiotic Therapy

Prophylaxis

Drug Regimen

Chemoprophylaxis for GBS

Tocolysis

Supplemental Progesterone

Hospitalization versus Home Care

Maternal Monitoring

Fetal Monitoring

- The optimal regimen is not clear, but a regimen with reasonable activity against major pelvic pathogens should be utilized [53].
- A seven day course of antibiotic prophylaxis is recommended for all women with PPROM who are managed expectantly.
- A current preference is: Ampicillin 2 gm intravenous every six hours for 48 hrs followed by Amoxicillin 500 mg po TID or 875 mg po BID for an additional five days.
- In addition, one dose of azithromycin 1 gm po is recommended upon diagnosing PPROM.
- Targets of antibiotic therapy involve:
 - Ampicillin and Amoxicillin for:
 - group B streptococcus
 - many aerobic gram negative bacilli
 - some anaerobes
 - Azithromycin for:
 - mycoplasma (an important cause of chorioamnionitis with PPROM)
 - Chlamydia trachomatis (cause of neonatal conjunctivitis and pneumonitis)
 - abnormal cerebral ultrasound prior to hospital discharge
- The regimen discussed is similar to the National Institute of Child Health and Human Development (NICHD) Maternal-Fetal Medicine Units (MFMU) Network trial.
 - This trial studied antibiotic therapy for reduction of infant morbidity after PPROM with IV ampicillin 2 g every 6 hours and erythromycin 250 mg every 6 hours for 48 hours FOLLOWED by oral amoxicillin 250 mg every 8 hours and erythromycin 333 mg every 8 hours for five days [16]. This is recommended by ACOG.



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- GBS chemoprophylaxis is indicated when GBS test results are positive or unknown and delivery is imminent. It is generally not given when women have had recent negative results, within the prior five weeks.
- For women GBS colonized, in labor at the time of admission or who go into labor within 48 hours of admission, IV portion of the regimen described (ampicillin 2 gm IV Q 6hr for 48 hours) should provide adequate treatment.
- The regimen of IV ampicillin followed by oral amoxicillin, combined with azithromycin, is usually given for seven days.
- Once the seven day regimen is complete, the antibiotics should be discontinued. For a woman with culture positive GBS, specific prophylaxis for GBS should be resumed when the patients goes into subsequent labor [55].
- If the woman has a history of high risk for penicillin allergy it is suggested she be given:
 - Azithromycin 1 gram orally upon admission, plus
 - Clindamycin 900 mg intravenously every 8 hours for 48 hours, plus
 - Gentamicin 5 mg/kg actual body weight intravenously every 24 hours for two doses, FOLLOWED BY
 - Clindamycin 300 mg orally every eight hours for five days

This regimen is appropriate for patients with a positive GBS culture and laboratory-documented GBS susceptibility to clindamycin
- If the women has a low-risk penicillin allergy antibiotic therapy suggested is a one time oral dose of azithromycin 1 g and cefazolin 1 g IV Q 8 hours for 48 hours, followed by cephalexin 500 mg orally four times daily for five days. These drugs provide coverage for both GBS and Escherichia coli, which are two major causes of neonatal infection.
- If the woman is at high risk for anaphylaxis and GBS is resistant to clindamycin per lab results the following treatment regimen is recommended:
 - Azithromycin 1 gram orally upon admission, PLUS
 - Vancomycin 20 mg/kg every 8 hours (maximum single dose 2 grams) for 48 hours
- [Algorithm's are available for GBS prevention from the Centers for Disease Control and Prevention](#)





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- Tocolysis has a principal indication in the setting of PPROM in order to delay delivery 48 hours allowing for administration of corticosteroids.
- Tocolysis should not be administered beyond 48 hours as a general rule.
- Tocolysis is not indicated when women are found to have advanced labor (> 4 cm dilated) or in those with findings to suggest the presence of subclinical or overt chorioamnionitis.
- Tocolysis is also contraindicated with non-reassuring fetal status, abruptio placentae, and significant risk of cord prolapse such as fetal malpresentation and dilated cervix.

Management



Click each term to learn more.



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- There is no evidence so suggest supplemental progesterone extends the latent period in women with PPROM or provide other benefits.
- However, women on supplemental progesterone due to a prior pregnancy with preterm delivery related to preterm labor or PPROM, discontinuing the progesterone upon diagnosis of PPROM would be appropriate.

Management



*Click each term to
learn more.*



**Administration of Antenatal
Corticosteroids****Antibiotic Therapy****Prophylaxis****Drug Regimen****Chemoprophylaxis for GBS****Tocolysis****Supplemental Progesterone****Hospitalization versus Home Care****Maternal Monitoring****Fetal Monitoring**

- Women with PPROM and a viable fetus warrant hospitalization from the time of diagnosis until delivery, with very few exceptions.
- Once PPROM is diagnosed, the woman's activity is limited to using the bathroom and sitting up a bedside chair.
- All pregnant women hospitalized and placed on bedrest should undergo thromboprophylaxis with sequential compression devices [56].
- Women with additional risk factors for deep venous thrombosis (DVT) should receive prophylactic doses of enoxaparin (Lovenox), 1 mg/kg/day. Enoxaparin (Lovenox) should be discontinued 48 hours prior to delivery.
- When weight gain occurs during pregnancy, intermediate dose anticoagulation adjustment can be made. American College of Physicians [75] and ACOG [76] recommend enoxaparin 40 mg twice daily, some experts use an alternate dose of enoxaparin at 1mg/kg.



Administration of Antenatal Corticosteroids

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- Women with PPROM need to be monitored for signs of infection. Unfortunately, there is not a consensus to the best approach for these women.
- Minimally, routine parameters such as maternal temperature, uterine tenderness and contractions, maternal and fetal heart rates, should be monitored.
- Markers for inflammation and infection may be periodically ordered but have not proven useful [57].
- It is controversial for the woman to undergo amniocentesis for Gram stain, culture, leukocyte esterase, and glucose.
- An amniocentesis is not recommended to screen for amniotic infection in an asymptomatic woman.
- The amniocentesis may be indicated when the clinical diagnosis of chorioamnionitis is uncertain and more information is needed for treatment versus expectant management.
- In at least 50 percent of patients, the amniotic fluid sample is insufficient for the woman to undergo amniocentesis, thus the diagnosis of chorioamnionitis will have to be based clinically with indirect testing such as an abnormal peripheral white blood cell count and physical exam.
- Diagnosis and management of intraamniotic infection is beyond the scope of this program.



**Administration of Antenatal
Corticosteroids****Antibiotic Therapy****Prophylaxis****Drug Regimen****Chemoprophylaxis for GBS****Tocolysis****Supplemental Progesterone****Hospitalization versus Home Care****Maternal Monitoring****Fetal Monitoring**

- Fetal surveillance is generally employed such as kick counts, NSTs, and biophysical profile (BPP). These provide the woman and clinician some assurance of fetal well being.
- Recommendations to perform a daily NST is reasonable.
- When the NST is non-reassuring, then a BPP is obtained.
- When attempting to predict fetal infection, these tests (NST, BPP) do not have good sensitivity with NST at 39 percent and BPP at 25 percent [58].
- The optimum type and frequency of testing does not have a consensus.
- A low amniotic fluid volume is associated with an increased risk of fetal umbilical cord compression and shorter latency, but the value of NST's or the BPP score for predication of adverse fetal/neonatal outcome is low [15].
- Fetal umbilical cord doppler surveillance, similarly, is not useful for predicting fetal status in PPROM either [59-61].



The fetus/neonate is at greater risk of PPROM-related morbidity and mortality than the mother.

- Neonatal sepsis can occur from intrauterine infection with long term neurodevelopmental abnormalities, particularly cerebral palsy.
- Umbilical cord compression can lead to fetal asphyxia.
- Limb restriction deformities and pulmonary hypoplasia may result from the oligohydramnios; primarily with severe oligohydramnios in early to mid second trimesters.
 - When membranes rupture occurs after 23 weeks, these complications are rare.
- Umbilical cord prolapse may lead to fetal asphyxia.
- Abruptio placentae may also lead to fetal asphyxia.
- Preterm birth may lead to:
 - Respiratory abnormalities
 - Intraventricular hemorrhage
 - Necrotizing enterocolitis
 - Retinopathy of prematurity
 - Patent ductus arteriosus





Click the arrows below to view the slides.

Women with HSV, HIV, or cerclage

- Controversy exists with the expectant management of women with PPRM and genital herpes simplex virus (HSV) or human immunodeficiency virus (HIV) and opinions regarding the best course of action diverge widely.
- These issues are beyond the scope of this program.



Slide 1 of 5





Click the arrows below to view the slides.

Meconium stained fluid

- Term and preterm women with PROM studies showed those with meconium stained amniotic fluid have higher rates of both overt and subclinical chorioamnionitis and positive amniotic fluid cultures [61-63].
- When meconium is released it predisposes to infection by enhancing growth of bacteria and lowering phagocytic capacity of neutrophils [64].
- It is possible, however, that the meconium-like staining could actually be pigment staining associated with decidual hemorrhage such as from abruption.
- Women with PPROM and meconium-stained amniotic fluid should be evaluated for signs of chorioamnionitis.
- Meconium in isolation is not an indication for intervention. So no action is needed in the absence of these signs.



Slide 2 of 5





Click the arrows below to view the slides.

Tissue sealants

- Neither the safety nor the efficacy has been established for tissue sealants. However, there are some case reports of a variety of tissue sealants (ie fibrin glue, gelatin sponge) showing some success in stopping leakage.



Slide 3 of 5





Click the arrows below to view the slides.

Amnioinfusion

- Pregnancy outcomes were studied in a 2014 systematic review and meta-analysis comparing women who received antepartum transabdominal amnioinfusion to those who underwent usual care for PPROM in the third trimester. The five trials had n = 241 pregnancy women.
- Very small trials with low to moderate quality have shown transabdominal amnioinfusion resulting in statistical reductions in neonatal death, sepsis, infection, and pulmonary hypoplasia but the data for each outcome were limited.
- Better understanding is needed on peri-natal outcome regarding whether amnioinfusion is beneficial in PPROM. More and better information is needed about the effects of specific amnioinfusion protocols:
 - Selection of patients such as gestational age at membrane rupture
 - Other interventions with type, dose, and duration of antibiotics along with use of corticosteroids
 - Amnioinfusion in women with PPROM is not indicated until further research is available to support a change in practice.



Slide 4 of 5





Click the arrows below to view the slides.

Diagnosis and treatment of overt infection

- When there is maternal fever with associated leukocytosis, maternal and fetal tachycardia, uterine tenderness, and malodorous discharge, overt chorioamnionitis can be clinically diagnosed.
- Subclinical chorioamnionitis diagnosis requires amniocentesis to identify microorganisms in the amniotic fluid with gram stain and culture along with documenting an abnormally low amniotic fluid glucose concentration.
- In some countries evaluation of infection can be performed using a rapid test for interleukin-6 (IL-6). This is the most sensitive marker for microbial invasion of the amniotic sac.
- Women who develop overt infection require therapy with therapeutic, rather than prophylactic, antibiotics.
- Of note, women with PPROM with an identifiable genital tract infection such as gonorrhea, chlamydia, bacterial vaginosis, that would not be eliminated with prophylactic antibiotic regimen should receive antibiotics specifically targeting the infection.

« Slide 5 of 5 »

- Magnesium sulfate for neuroprotection:
 - Magnesium sulfate is given prior to delivery according to standard clinical protocols for fetal neuroprotection in women at gestational ages of at least 24 weeks but less than 32 weeks at risk of imminent delivery.
- Timing of delivery for expectantly managed pregnancies:
 - The approach to timing delivery is based on weeks gestation.





Gestation at 24 - 32 Weeks



Gestation at 32w0d - 33w6d



Gestation at 34 Weeks+



Click the labels above to learn about delivery at each stage.



Gestation at 24 - 32 Weeks



Gestation at 32w0d - 33w6d



Gestation at 34 Weeks+

24 - 32 Weeks

- Plan
- When there is no evidence of chorioamnionitis or fetal compromise, then expectant management is appropriate.
- Administer corticosteroids
- Antibiotics, broad spectrum, are given and GBS prophylaxis at delivery, if indicated
- Generally, prematurity is the greatest risk to the fetus with uncomplicated PPRM at less than 34 weeks of gestation.
- It is appropriate to manage pregnancies at this gestational age expectantly in the absence of complications:
- Infection
- Abruption
- Cord prolapse
- Unstable fetal presentation
- Nonreassuring fetal assessment
- If delivery becomes evident, consideration to start magnesium for neuroprotection may be warranted.



Gestation at 24 - 32 Weeks



Gestation at 34 Weeks+

Gestation at 32w0d - 33w6d

32w0d - 33w6d

- Plan
- When fetal lung maturity can be documented or there is evidence of an intraamniotic infection clinically or on amniocentesis, then delivery is indicated.
- Otherwise expectant management with delivery at 34 weeks
- Administer corticosteroids if there is evidence of fetal lung immaturity or fetal lung status is unknown
- Administer broad spectrum antibiotics and GBS prophylaxis, if indicated



Gestation at 24 - 32 Weeks



Gestation at 32w0d - 33w6d



Gestation at 34 Weeks+

34 Weeks +

- When the pregnancy reaches 34 weeks, usually labor is induced without testing for fetal lung maturity.
- However, if there is uncertainty about the patient's gestational age, attempting to aspirate amniotic fluid from the vaginal vault to test for fetal lung maturity, is reasonable.
- Use of the lamellar body count as the initial screen for fetal lung maturity is appropriate.
- If this test is immature, performing a lecithin/sphingomyelin (L/S) ratio is a common next step.
- When testing shows a low risk of neonatal respiratory problems, initiating delivery is appropriate because the risks of prematurity are small when compared to the risk of developing maternal or fetal complications during expectant management [67].
- When fluid can not be aspirated or testing suggests a high risk of neonatal respiratory problems, continuing management expectantly until 36 weeks (by estimate) is appropriate; delivery without resampling at this time is appropriate.
- The woman who develops clinical evidence of infection or abruption, preterm labor or non-reassuring fetal well being would be indication for earlier delivery
- In summary the plan for gestations 34wks 0days or greater is:
- Delivery
- Corticosteroids are not indicated
- GBS prophylaxis begun on admission and continued until delivery unless she is known to have negative GBS status.

- In the situation where there are no contraindications to labor and vaginal birth, most women will deliver by spontaneous or induced vaginal delivery [68].
- Cesarean delivery is performed for standard indications, otherwise, labor is induced.
- At this point, performing a digital cervical exam is indicated to determine whether cervical ripening has occurred.
- Oxytocin is administered for induction when the cervix is found to be favorable, according to standard protocols.
- Once cervical ripening is adequate, the use of oxytocin over prostaglandins is favored since oxytocin is more easily titrated.



Unfavorable cervix

- With an unfavorable cervix, it may be advantageous to utilize misoprostol since it is effective for inducing labor as well.
- Fifteen randomized trials of women with term PROM underwent a meta-analysis and reported the rate of vaginal delivery in 12 and 24 hrs was similar after administration of misoprostol or oxytocin [69].
- Is it unknown whether misoprostol was advantageous in the subgroup of women with an unfavorable cervix since it was not evaluated.
- The optimum dose and route of misoprostol administration have not been determined.
- Prostaglandin E2 is a reasonable alternative [70].
- There is minimal information on the safety of mechanical methods of cervical ripening in PROM [71].
- Some methods are not utilized due to concerns of introduction of a foreign body may increase infection, such as Foley bulb for cervical ripening; so this method is typically avoided.



*Click each box for
more information.*





PPROM refers to rupture of fetal membranes prior to labor in pregnancies < 37 weeks 0 days.

PPROM is responsible for one-third of preterm births and occurs in three percent of pregnancies.

- PPROM risk increases three fold when PPROM has occurred in a prior pregnancy.
- PPROM is a clinical diagnosis based on visualizing amniotic fluid in the vagina of a women presenting with a history of leaking fluid.
- PPROM, by clinical history, should be confirmed by visualizing or diagnostic testing while excluding other causes of wetness:
 - Urinary incontinence
 - Vaginal discharge
 - Perspiration



Click each box for more information.





A clinical history suggestive of PPROM should be confirmed by visual inspection or diagnostic testing to exclude other causes of wetness, such as urinary incontinence, vaginal discharge, and perspiration.

Managing a women with PPROM is based on several factors:

- Gestational age.
- Availability of neonatal intensive care
- Presence or absence of maternal or fetal infection
- Presence or absence of labor or abruptio placentae
- Stability of fetal presentation and fetal heart rate tracing pattern
- Probability of fetal lung maturity
- Cervical status

Expeditious delivery of women with PPROM is clinically appropriate if intrauterine infection, abruptio placentae, nonreassuring fetal testing, or a high risk of cord prolapse is present or suspected.



**Click each box for
more information.**





Expeditious delivery of women with PPROM is clinically appropriate if intrauterine infection, abruptio placentae, nonreassuring fetal testing, or a high risk of cord prolapse is present or suspected.

Women with PPROM must be expeditiously delivered if any of the following occur:

- Intrauterine infection
- Abruptio placentae
- Nonreassuring fetal status
- High risk of cord prolapse



Click each box for more information.





For stable patients with PPROM < 34 weeks, expectant management is appropriate. In addition:

- Administer a course of antenatal corticosteroids to enhance fetal lung maturation in pregnancies less than 34 weeks.
- Prophylactic antibiotics with:
- Ampicillin 2 g IV every 6 hours and erythromycin 250 mg every 6 hours for 48 hours FOLLOWED by oral amoxicillin 250 mg every 8 hours and erythromycin 333 mg every 8 hours for five days
- Azithromycin po one gram once at the time of admission and repeat the dose five days later
- For women with a penicillin allergy recommended antibiotic therapy is a one time oral dose of azithromycin 1 g and cefazolin 1 g IV Q 8 hours for 48 hours, followed by cephalexin 500 mg orally four times daily for five days.



**Click each box for
more information.**

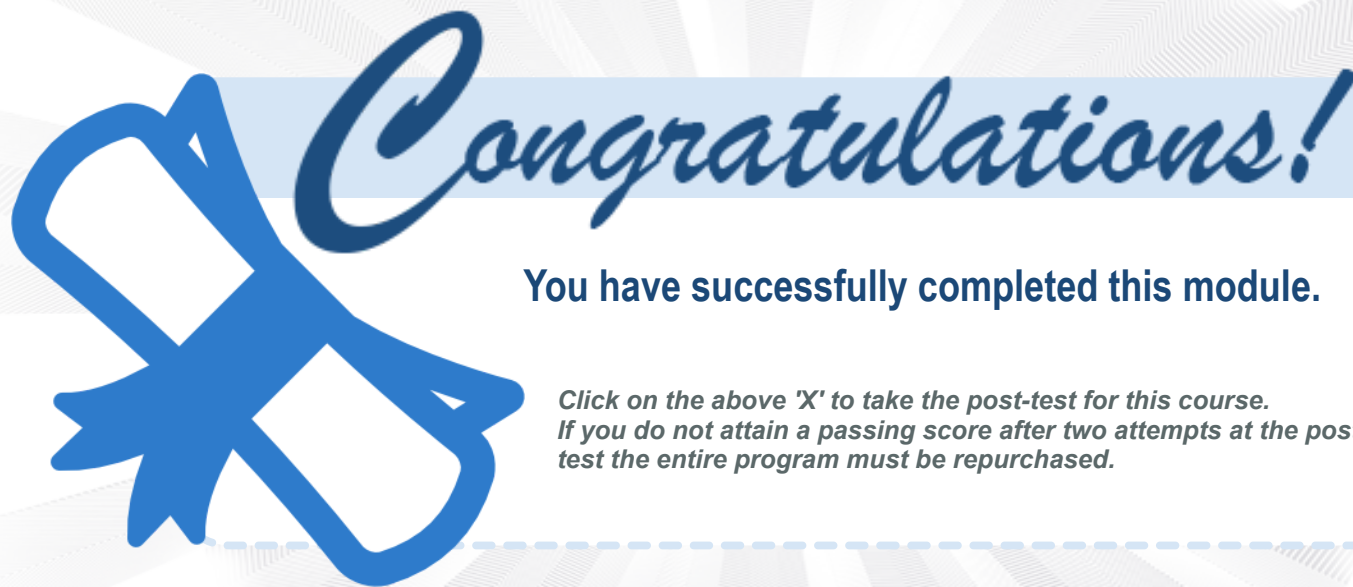




Women with confirmed gestational age, delivery at 34 weeks or beyond without assessing pulmonary maturity is appropriate.

If gestational age is uncertain, attempting to confirm lung maturity before delivery is appropriate.

Assuming the mother and fetus are stable, then delivery at 36 weeks is reasonable when the amniotic fluid cannot be obtained or the test demonstrates lung immaturity.



Congratulations!

You have successfully completed this module.

*Click on the above 'X' to take the post-test for this course.
If you do not attain a passing score after two attempts at the post-test the entire program must be repurchased.*

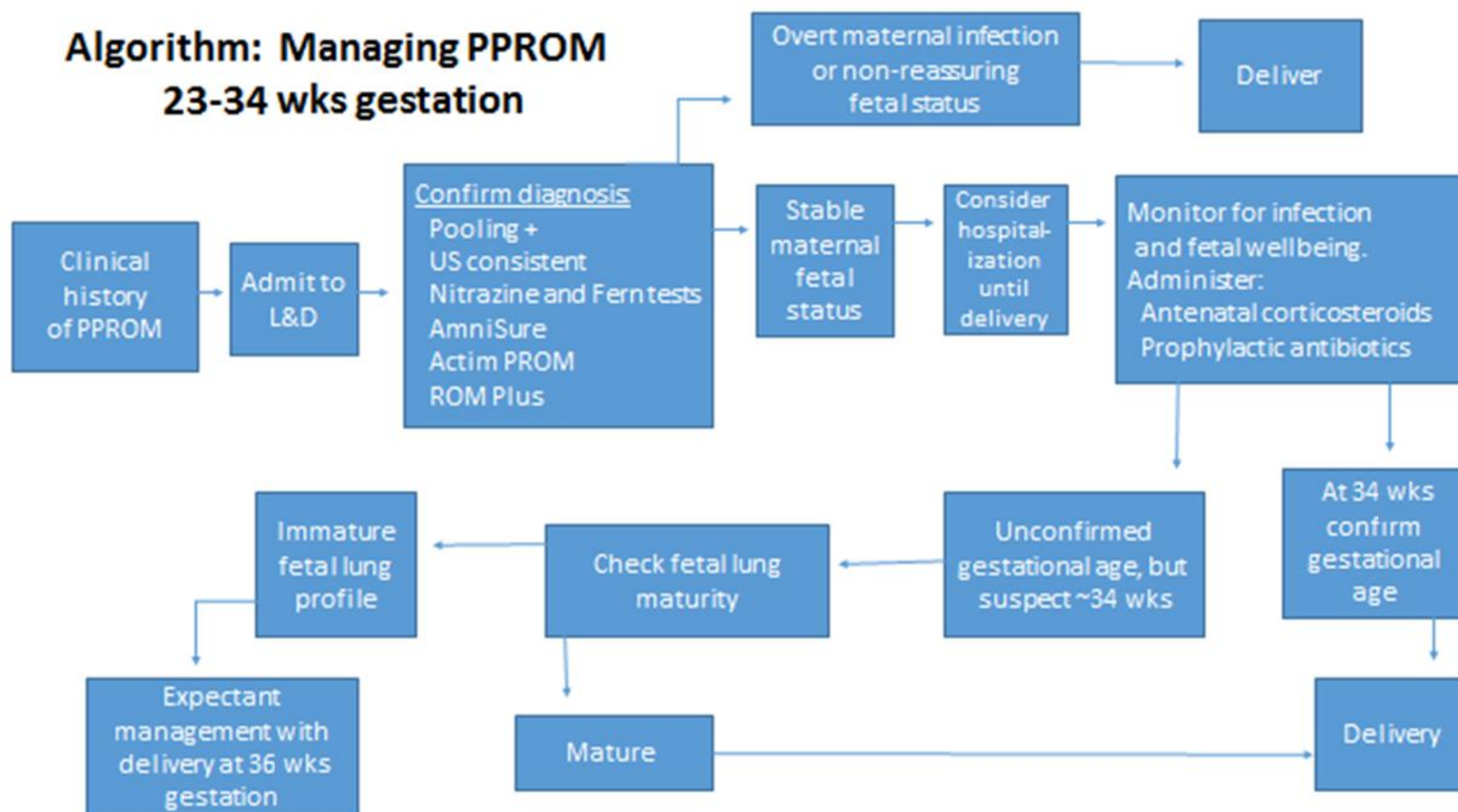
Risk Factors Table

Bacteruria	No partner
Periodontal disease	Low socioeconomic status
Placenta previa	Anxiety
Placental abruption	Depression
Vaginal bleeding, especially in more than one trimester	Life events, divorce, separation, death
Previous PTB	Abdominal surgery during pregnancy
Substance abuse	Occupational issues
Smoking	upright postures
Maternal age (younger than 18 or older than 40)	use of industrial machines
African-American race	physical exertion
Poor nutrition and low body mass index	mental or environmental stress related work
Inadequate prenatal care	working conditions
Anemia (hemoglobin <10g/dL)	Multiple gestation
Excessive uterine contractions	Polyhydramnios
Low level of educational achievement	Uterine anomaly
Genotype	DES induced uterine changes
Fetal anomaly	History of second trimester abortion
Fetal growth restriction	History of cervical surgery
Environmental factors (i.e. heat, air pollution)	Premature cervical dilatation or effacement (shortened cervix)
STI's	

Table 2: Complications with PPROM

Pregnancy Complication	Potential Consequences for Offspring	Potential Maternal Consequences
Intrauterine infection	Neonatal sepsis Long-term neurodevelopmental abnormalities, particularly cerebral palsy	Postpartum endometritis
Umbilical cord compression	Fetal asphyxia	Cesarean delivery
Oligohydramnios	Limb restriction deformities and pulmonary hypoplasia (primarily with severe oligohydramnios in the early to mid second trimester), occurring rarely when membranes rupture occur after 23 weeks.	
Fetal malpresentation		Cesarean delivery
Umbilical cord prolapse	Fetal asphyxia	Cesarean delivery
Placental abruption	Fetal asphyxia	Cesarean delivery Coagulopathy
Preterm Birth	Morbidity of prematurity	

Algorithm: Managing PPROM 23-34 wks gestation



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