MERITER IRB #: 2020-003

RESEARCH PROTOCOL TITLE: PROMMO Trial: Prelabor rupture of membranes managed with oral misoprostol versus intravenous oxytocin

PRINCIPAL INVESTIGATOR: Jacquelyn H Adams, MD

Clinical Trials.Gov Listing at:
https://clinicaltrials.gov/ct2/show/NCT04478942?term=NCT04478942&draw=2&rank=1

STUDY PURPOSE

The purpose of this study is to look at labor induction for women who have broken their water but aren't yet in labor and have a cervix that is not considered favorable for labor.

RESEARCH PROCEDURES

Study participants will receive one of two medications as part of standard medical care. Participants will complete two surveys, which will take approximately 20 minutes.

ELIGIBILITY CRITERIA FOR SUBJECTS

1. Early Term to late term pregnancy (>37 weeks and 0 days and <42 weeks and 0 days)
2. Late Preterm Pregnancy (34 weeks and 0 days and <37 weeks)
3. Confirmed rupture of membranes by either sterile speculum exam or AmniSure
4. Simplified Bishop Score ≤ 6
5. Singleton gestation in women over 18 years old
6. Appropriate gestational age dating by certain LMP or ultrasound performed prior to 20 weeks gestational age

COMPENSATION/REIMBURSEMENT

No compensation is available for participation in this study.

RESEARCH LOCATION

UnityPoint Health – Meriter Hospital
Madison, WI 53715

CONTACT PERSON Jackie Adams, MD

PHONE: 608-417-6099