

# Low Dose Antenatal Corticosteroids for Late Preterm Delivery

NCT05698966

---

<b>Status</b>	RECRUITING
<b>Phase</b>	Not Applicable
<b>Sponsor</b>	Rambam Health Care Campus
<b>Enrollment</b>	1,510 participants

## Key Eligibility Criteria

---

### Inclusion (5)

- Criteria for determination of late preterm delivery risk:
- Preterm uterine contractions with intact membranes, and at least 3 cm dilation or 75% cervical effacement
- Spontaneous rupture of the membranes
- Expected preterm delivery for any other indication via induction or cesarean between 24 hours to 7 days after the planned randomization, as determined by the obstetric provider.
- \-

### Exclusion (5)

- Expected delivery in less than 12 hours, irrespective of cause including: 1) ruptured membranes in the presence of more than 6 contractions per hour or cervical dilation of 3 centimeters or more unless oxytocin was withheld for at least 12 hours (although other induction agents were allowed), 2) chorioamnionitis, 3) cervical dilation of 8 cm or more, and 4) evidence of non-reassuring fetal status requiring immediate delivery.
- Prior ACS treatment
- Current known or suspected infection ( viral, bacterial or other)
- Pre-gestational diabetes mellitus.
- Any infection that required antibiotics or hospitalization in the month prior to study allocation - Poor understanding of the informed consent language

## Locations (16 total)

---

Emek Medical Center, Afula, Israel  
Kaplan Medical Center, Ashkelon, Israel  
Soroka Medical Center, Beersheba, Israel  
... and 13 more locations

---

<https://clinicaltrials.gov/study/NCT05698966>

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at [ClinicalTrials.gov](https://clinicaltrials.gov). Generated by [ClinicalTrialsFinder.org](https://clinicaltrialsfinder.org).