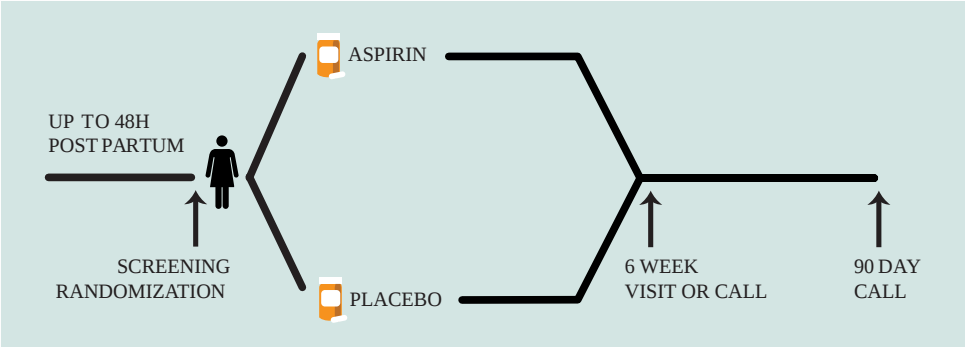
Pilot **PARTUM** Trial:

Postpartum **A**spirin to **R**educe **T**hromboembolism **U**ndue **M**orbidity

Postpartum women at modest risk of VTE (deep vein thrombosis or pulmonary embolism) will be randomized to low-dose aspirin daily versus placebo daily for 6 weeks postpartum. Participation is simple:



Thank you for supporting the pilot PARTUM trial!

partumtrial.ca

OMNI Research Group Pager:
1-855-266-7243 then code 666474#

INCLUSION

ONE of:

1. Mild inherited thrombophilia:
Heterozygous FVL or Heterozygous PGM or PC def or PS def

2. Bedrest (90%) for ≥7 days during pregnancy

OR

TWO of:

3. Pre-pregnancy BMI ≥30 kg/m^2

4. Smoking in past 12 months

5. Past superficial vein thrombosis

6. Pre-eclampsia

7. Stillbirth in current pregnancy (>20 weeks)

8. SGA infant <3rd percentile

9. Unplanned cesarean delivery

10. Postpartum infection

11. Postpartum hemorrhage

PARTUM

Subjects are excluded if they are >48 hrs after delivery, received >2 LMWH doses or have had a prior VTE.
See poster for full eligibility criteria.