Partum: A pilot study assessing the feasibility of a randomized controlled trial evaluating aspirin in postpartum women at risk of developing venous thromboembolism. Protocol Number REB19-1237 Version V1.5 08Jun2020

OHSN Protocol ID # 20200832

Participant Name: _____ MRN: _____

INFORMED CONSENT PROCESS

VISIT DATE ___ ___ /___ ___ ___ /___ ___ ___ ___ / (D/M/Y)

1) Has the patient signed the main study consent form? Yes □ No □

Date consent signed ___ ___ /___ ___ ___ /___ ___ ___ ___ / @ _____: _____ hrs.

Consent Process for Study Title

The details of the above study were discussed with the participant. The study was explained in detail including all the contents of the informed consent document. The participant was encouraged to ask questions. All questions were answered to the satisfaction of the participant. The participant was given adequate time to read the informed consent form **(ICF) version _____** and the opportunity to discuss it. **ICF version _____ was** signed without alteration by the participant.

The original copy of the signed informed consent document was placed in secure study file and a photocopy given to the participant.

Consent obtained by _____ Signature: _____

Date: __ __ / __ __ __ / __ __ __ __ /

Addendum

_____
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