WORKFLOW EXAMPLE 1:

Notes:

- Pharmacy timelines between 8am-5pm M-F not weekends, we can confirm with pharmacy.
- JHS Nurses certain shifts of availability- when study team needs their assistance. Please inform weekdays and hours, if possible.
- The workflow can also be color coded i.e blue for JHS and orange for UM or just differentiate one institution from the other.
- -Location is key.

Inpatient	Inpatient
Screening for eligibility	Attending Neurologist Dr. X – PI –ER (JHS)
Informed Consent process	Study Coordinator (name) – ER (JHS)
Review of the SOC Brain qualifying Images,SOC labs, Data abstraction	Study Coordinator (name) – EB(JHS)
Randomization	Study team — (name if possible) (JHS)
Administration of Investigational Product: IV INFUSION (Please provide details to the Nursing team i.e. 2ml every 3hrs, duration of the infusion etc)	JHS NURSE - JHS ANGIO SUITE
Standard of Care Endovascular treatment	Neuro Intervention Dr. (റുമ്പ്റ്റു if possible), Sub-PI /JHS Angio Suit
Day 1 post EVT-Review medical record to check the for adverse events. Perform study-assessments	Study team/Study Coordinator (name) – ICU (JHS)
Days 2-6 post EVT or at discharge if sooner-Review medical record to check the for adverse events. Review SoC brain imaging. Perform study assessments	Study team/Study Coordinator (name) – ICU / WW 15 (JH
Outpatient	Outpatient
Vist 5 (Day 30-Review medical record to check the for adverse events. Review SoC brain imaging. Perform study-assessments)	ALL OUTPATIENT ACTIVITIES AND FOLLOW UP VISIT WILL BE PERFORMED AT UM BY STUDY TEAM
Visit 4 Day (Day 90- Review medical record to check the for adverse events. Review SoC brain imaging. Perform study-assessments	

WORKFLOW EXAMPLE 2:

a prospective cohort study designed to evaluate the longitudinal health outcomes of young women

	. Only Pregnant women will be enrolled at JHS.
2.	Study coordinators (SC) will be in constant communication with the OB department. They will attend the weekly OB meeting where patients that are coming in for care are presented. In this meeting, the study coordinator will learn of any possible candidate to be approached for the study.
3.	is the treating physician for these patients, and she is part of the study. and JHS registered nurses will know in advance about the study and will help the study team to identify possible candidates. Upon identification of a potentially eligible pregnant woman, the PCP will introduce the study to the patient. For women who express interest in learning more about the study, the PCP will introduce them to the study team to provide a full explanation of the study, education, and counseling as part of the study informed consent process.
4.	 Study Workflow: Recruitment and informed consent will take place at clinic ACC West Building by the study coordinator Specimen collection, interviews, and assessments for the pregnancy visit will occur at UM Batchelor's Children's Research Institute (BCRI) The delivery visit will take place at Jackson (Holtz), JHS nurses will help the SC in the collection of repository specimens (serum, plasma), maternal hair, maternal swabs (vaginal), and they will be processed and shipped out centrally by UM. If the delivery visit cannot be completed at JHS, it will be scheduled to occur within two weeks of delivery at Batchelor's Building.
5.	All the rest of the visits and the non-pregnant cohort will occur at UM Batchelor's Building.

6. Chart abstraction will be performed annually using data available on Cerner and Epic for both groups of

participants.