Research Study
A Prospective, Double Blind, Randomized, Placebo Controlled Study of IV ibuprofen and IV acetaminophen Post Uterine Fibroid Embolization Procedure for the Control of Pain and Nausea

IRB Number: 14-000359

Study Information
If you are between 21 – 60 years of age, you may be eligible to participate in a research study to collect information related to the use of certain medications given during and after a UFE procedure for the treatment of pain and nausea. This information is expected to help your doctors better manage the pain and nausea associated with a UFE procedure.

Study Procedures
- Pain and nausea assessments at the completion of the UFE procedure, 6 hours after the procedure and at the time you are discharged from the hospital.
- A Patient Satisfaction Questionnaire 2 week after the UFE procedure

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