**The Cleveland Clinic Foundation**
**Information Sheet to Participate in a Research Study**

# Current practice strategies for post procedure analgesia after percutaneous needle tenotomy

**Principal Investigator:** Michael Dakkak, DO (954) 659-5430

## What is the purpose, procedures and duration of this study?

We invite you to take part in a research study because you may perform tenotomy procedures, a procedure to treat pain in tendon(s). The purpose of this study is to evaluate current pain management prescribing practices after percutaneous needle tenotomy. This research will be the first to investigate post procedural analgesia management practices after this procedure.

You will be asked to take a brief 5-10 minute survey; this research involves analyzing prescribing practices after percutaneous needle tenotomy.

## What are the risks of participating in the research study?

### Confidentiality Risks

There is a potential risk of loss of confidentiality of your data. Every effort will be made to keep your information confidential through the use of the following safeguards: safeguards which reduce the use of PHI when feasible, limit data access to the study personnel and storing data in secured manner are in place, anonymous surveys in which will not be linked back to you.

### Questionnaire/Survey Research

Some of the questions we will ask you as part of this study may make you feel uncomfortable. You may refuse to answer any of the questions and you may take a break at any time during the study. You may stop your participation in this study at any time.

## What are possible benefits of participating in the research?

There is no personal benefit to you by participating in this research study. The knowledge to be gained from this research may be beneficial for other patients, society or science.

Your participation is completely voluntary. If you have any questions regarding the study, contact the Principal Investigator, Michael Dakkak at (954) 659-5430. If you have any questions regarding your rights as a participant, contact the Institutional Review Board at (216) 444-2924.

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