SHORT TITLE: Maverick-HCM

RESEARCH PROTOCOL TITLE: A Randomized, Double-blind, Placebo-controlled, Concentration-guided, Exploratory Study of Mavacamten (MYK-461) in Patients with Symptomatic Non-obstructive Hypertrophic Cardiomypathy

PRINCIPAL INVESTIGATOR: John Moses, MD

RESEARCH SPONSOR: MyoKardia, Inc.

Clinical Trials.Gov Listing at:
https://clinicaltrials.gov/ct2/show/NCT03442764?term=mavacamten&cond=Hypertrophic+Cardiomyopathy&rank=3

STUDY PURPOSE
This study evaluates the safety and tolerability of a 16-week course of mavacamten (the investigational medication) in patients with symptomatic Hypertropic Cardiomyopathy (a condition where the heart becomes thickened without an obvious cause).

STUDY PROCEDURES
Participation in the study can last up to 28 weeks with 8 in clinic visits and three phone call visits. During clinic visits blood will be drawn, an ultrasound of the heart may be done, a treadmill test may be done, questionnaires may be done and a physical exam will be done.

ELIGIBILITY CRITERIA FOR SUBJECTS
1. 18 years of age or older
2. Be able to walk on a treadmill
3. Give informed consent and comply with study procedures

SUBJECT PAYMENT
Compensation will range from $25-$75 per visit.

RESEARCH LOCATION AND CONTACT PERSON
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