

## **STUDY OVERVIEW**

Mind Star Research is conducting an observational study to document how Pharmacogenomics results affect patients' plan of care and changes in medication prescription(s).

We would like to invite you to participate in this study as a Principal Investigator. This is NOT a clinical trial or a double-blind study.

## **OBSERVATIONAL STUDY PROTOCOL**

Medical Provider Experience and Assessment on Documenting Prescribing and Plan of Care Changes Due to Pharmacogenomics Testing.

Protocol Number: PAS1463

Initial Version Date: March 22, 2017

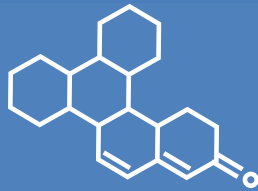
Study product: Observational Study

## **PRINCIPAL INVESTIGATOR CRITERIA**

- Must be a Medical Practitioner:
- Medical Doctor (MD) • Doctor of Osteopathic (DO) • Physician Assistant (PA)
- Advanced Practice Registered Nurse (APRN) • Nurse Practitioner (NP)

## **DATA POINT COLLECTION CRITERIA**

- Patients with medical necessity
- Male or Female, ages 18 – 64
- Must have insurance coverage for laboratory testing
- No Federally funded insurance plans



## STUDY COMPENSATION AND RECOGNITION

- 48 States \*Excluding California and New York
- Study length 36 months
- Compensation based on Third Party Fair Market Value assessment
- All Principal investigators are invited to be referenced in the Research Journals
- Study is registered with Institutional Review Boards (IRB)
- Study is pending acceptance to [www.clinicaltrials.gov](http://www.clinicaltrials.gov)

## CLINICALTRIALS.GOV

A service of the U.S. National Institutes of Health

ClinicalTrials.gov is a registry and results database of publicly and privately supported clinical studies of human participants conducted around the world.

Currently lists studies with locations in 50 States and 193 Countries

## THE FORMAT OF THE STUDY IS SIMPLE:

**1 ENROLL AS A PRINCIPAL INVESTIGATOR**

We are available 24/7 to answer any questions.

**2 SUBMIT A DAILY /WEEKLY SOURCE DOCUMENT**

Fill out Daily/weekly Source Document and fax to Mind Star Research.

**3 SUBMIT QUARTERLY CASE REPORT FORM**

Document the previous quarters' data collection.

**4 REPEAT**

Repeat until you receive protocols for the next phase.

I am here to help! Please let me know if you have any questions or if you are ready to enroll as a Principal Investigator. We look forward to working with you!