Reduction of Adverse Drug Events and Readmissions

NCT02738047

Status	RECRUITING
Sponsor	ClinLogic LLC
Enrollment	280,000 participants

Key Eligibility Criteria

Inclusion (4)

- Male or female patients of 25 years of age or older who are able to give their written Informed Consent to participate in a Clinical Study based on voluntary agreement with a thorough explanation of the patient's participation will be provided to them.
- Patient underwent PGx testing for alleles appropriate to the target drugs within the prior 120 days ("index PGx test assessment");
- Patient was receiving at least one medication known to be associated with allelic variation at the time of the ("index PGx test assessment"), including over-the- counter medications;
- Patient has a history of at least one TDAE over the 24-month period preceding the PGx test assessment, or has experienced inadequate efficacy from a target drug.

Exclusion (5)

- Patients will be excluded from the Study if any of the following criteria apply:
- Patient is currently hospitalized;
- Patient's medical and medication history is unavailable over the 120-day period preceding the PGx test assessment;
- Patient is unable to provide an accurate history due to mental Incapacity;
- Patient is known to have undergone prior PGx testing for genes specific to the target drug(s), exclusive of the PGx test relating to this Study.

Locations (1 total)

MD@Home, York, Pennsylvania, United States