

In the Electronic Health Record Era, Do We Still Need Clinical Registries

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Duke Clinical Research Institute

FROM THOUGHT LEADERSHIP
TO CLINICAL PRACTICE

The Panel

- Lisa Wruck, PhD (moderator)
 - Director, Center for Predictive Medicine, Duke Clinical Research Institute
- Sharon Hensley-Alford, PhD
 - Associate Chief Health Officer at IBM Watson Health
- Matt Roe, MD, MHS
 - Professor of Medicine, Duke University Division of Cardiovascular Medicine
- Marc Berger, MD
 - Chair of Real World Evidence Product Advisory Board, Shyft Analytics



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What do we mean by “Clinical Registry”?

A patient registry is an organized system that uses observational study methods to collect uniform data (clinical and other) to evaluate specified outcomes for a population defined by a particular disease, condition, or exposure, and that serves a predetermined scientific, clinical, or policy purpose(s).

- Gliklich RE, Dreyer NA: Registries for Evaluating Patient Registries: A User’s Guide: AHRQ publication No. 07-EHC001. Rockville, MD. April 2007