

Web-based patient registry for clinicians and researchers, allowing efficient data collection for improved management of the global epileptic population.



- ★ Dashboard view for quick overview of a Clinician's assigned patients
- ★ Detailed patient summary
- ★ Multiple sophisticated data collection forms
- ★ Data entry approval system with levels of accreditation
- ★ Multiple registries for handling different patient sets
- ★ Automated scheduled follow up reminders
- ★ Condition comparison charting
- ★ Data extract
- ★ Reporting on key data
- ★ Encourages optimal management of the patient
- ★ Provides a central repository, accessible worldwide
- ★ Created in partnership with the Auckland District Health Board

A registry for capturing data on patient episodes triggered by a medical condition; enables tracking and monitoring of outcomes based on approved treatments provided.

Benefits of EpiNet

- ✦ Improve clinical care of patients
- ✦ Dynamic clinical support, designed to capture relevant clinical data only
- ✦ Control access and entry of data by role:
 - Read-only access
 - Research Assistant – requires approval from investigator
 - Investigator
 - Primary Investigator (owner) for a specific patient
 - Anonymised administrator access
- ✦ The investigator who enters the information (the primary investigator) determines who can access the record
- ✦ Low-cost, collaborative approach to research
- ✦ Accessible from anywhere in the world
- ✦ Information on patients with specific syndromes allowing for focused studies
- ✦ Identification of relatively large groups of patients with uncommon or rare syndromes, so that research in these patient groups can be facilitated
- ✦ Data collection tool, not intended for use as a clinical trial

