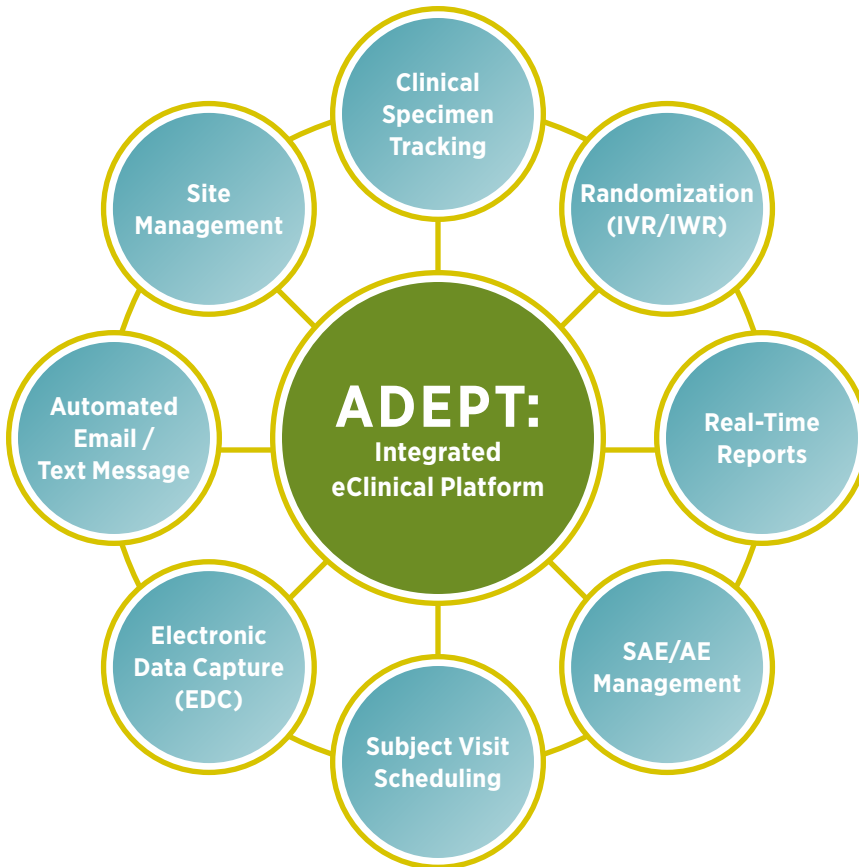


ADEPT: Integrated eClinical Platform

ADEPT greatly simplifies data collection via web-based Electronic Data Capture, while providing a comprehensive and integrated platform for the management of sites, clinical supplies, clinical specimens/images, and adverse events. All data entered by clinical sites is immediately available in the central database enabling sponsors to view standard and customized reports in real-time.



“ADEPT is NERI’s proprietary, web-based eClinical platform. It provides clinical trial sponsors with a dynamic, customizable tool for managing all of a study’s discrete, but connected, data collection functions through one integrated technology platform.”

**Patti L. Arsenault, CCDM,
PMP, MBA**

Director, Data Management and
Clinical Data Strategies

Although NERI has the capability of working with data from many commercially available data management systems, we offer ADEPT to sponsors with no license fees. Sponsors thereby benefit from dynamic, transparent, and cost-effective clinical study management.

ADEPT Advantages

- Fast customization and deployment
- Web-based data entry (no IT footprint)
- Native integration of common eClinical functions (EDC, IVR/IWR, supply tracking, document tracking, etc.)
- Integrated standard and customized reporting
- Adverse event alerts
- Real-time data management and analysis
- Extensive support and training
- Data security and integrity
- Proven utility and value

CASE STUDY: Global Congestive Heart Failure

Key Highlights:

- Phase III, randomized, double-blinded, placebo-controlled study in patients with Congestive Heart Failure
- 4,500 subjects
- 220 clinical study sites
- 6 countries (U.S., Eastern Europe, South America)
- Complex drug titration schedule
- Multiple lots of blinded drug/placebo

Details:

- Drug inventory management and distribution across all 12 drug/placebo codes to 220 sites, 4,500 subjects, across 6 countries (less expensive than traditional supply management)
- Investigational drug product tracking by lot number to each randomized subject
- Site drug supply needs anticipated based on subject allocation and drug expiry dates
- Drug and subject blinding and randomization (IVR/IWR) management
- Biologic specimen chain of custody tracking and reporting
- Site payment management based on milestones achieved
- Adverse event and study outcomes management
- Detailed subject and site management reports

To learn more about ADEPT,
please contact our Business Development Department:

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