EVADO Server

Design, execute and manage multiple trials and studies in a single web environment with Evado Server.

KEY FEATURES

- Built to exacting international quality standards.
- Capable of handling multiple clinical trials simultaneously that can be run across multiple trial sites.
- Can be accessed via the web on a desktop, an iPad or other mobile device.
- Integrates data collection processes.

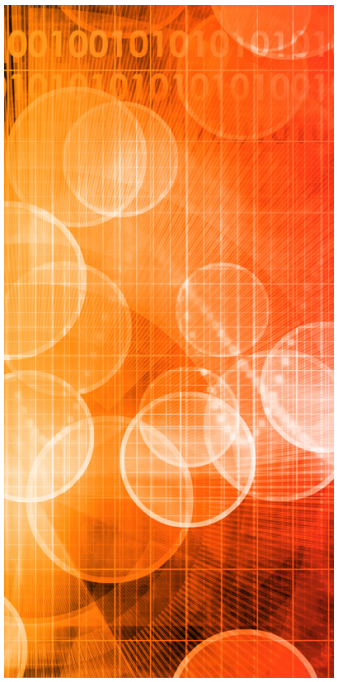
KEY BENEFITS

- Streamlines the data collection process.
- Saves time.
- Increases accuracy.
- Highly intuitive — easy to learn and use.

Evado Server provides a single web environment for the design, execution and management of multiple trials and studies.

Hosted securely in the cloud or on premises for larger institutions, Evado Server has been built to exacting international quality standards and meets the requirements of GCP ICH 6 and FDA Part 11. The Evado Server provides the core features and functionality for all of the Evado eClinical software modules including the EDC Clinical module that provides a web-based Electronic Data Collection environment and the Management or CTMS module that provides a web-based Clinical Trial Management environment for the execution of clinical trials.

The Evado Server has the following features:



- **General:**
  - Capable of handling multiple clinical trials simultaneously that can be run across multiple trial sites
  - User friendly, highly intuitive and easy to learn.
  - Designed to save time and costs by minimizing the site data collection time by pre-filling forms where appropriate, and then asking users to confirm that standard activities have been undertaken.
  - Integrates data collection processes.
  - Provides the capability to optimise the site navigation to streamline the data collection process.
    - The researcher confirms the trial activities that have been undertaken.
    - The site monitor confirms that the activities have been completed — this event triggers the invoicing process.
- **Centralised trial management environment**:
  - Evado can be accessed via the web on a desktop, an iPad or other mobile device.
  - Register subjects for one or more studies.
  - Initialise, start and close studies.
  - Select and manage trial subjects.
  - Select organizations involved in the trial.
  - Arrange visits with subjects.

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- Review trial data.
- The Principal Investigator is able to delegate authority to site staff and site coordinators.
- Handle multiple trial arms.
- Define visit schedules for each arm in the trial.
- Define and track one or more visit activities.

- **Trial record environment**:
  - Site staff can only access the subject data associated with the trial they are currently running.
  - Site staff can update the subject demographics online.
  - Site staff can generate patient-specific trial schedules of attendance.
  - Site staff can record the patient's attendance and record protocol visit compliance.
- **The administration environment only has to be set up once and can be used across multiple trials to define:**
  - User profiles
  - Global roles
  - Trial-specific roles
  - Site profiles.
- **FDA and GCP compliance**:
  - Control of user access to software components and functions.
  - Logging of user activity.
  - Creation and management of audit trails to track changes to all database records in accordance with FDA Part 11 requirements.

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