Evado Clinical

Evado Clinical is an award-winning cloud-based platform that manages clinical data for clinical trials, studies, and registries. Evado Clinical is Australian, developed, hosted, and supported.

Evado Clinical meets the regulatory requirements of FDA Part 11, HIPPA and the Code of Good Clinical Practice 6.

Evado Clinical provides a fully integrated clinical trial platform, allowing customers to run multiple trials. studies or registries across multiple sites.

Benefits

- Quick and easy to learn.
- Trials and studies can be quickly deployed.
- Simplifies clinical data management.
- Easy creation and completion of form records.
- Record layouts can be used across multiple trials, studies, and registries.
- Transparent pricing.
- fully integrated environment.

Key features

- Run an unlimited number of trials and studies simultaneously in the one environment.
- Online form tools for data collection.
- Online trial and study data management.
- Scheduling for multiple arm trials.
- Integrated Informed Consent for trials and registries.
- Integrated Clinical Outcome Assessments.
- Data exception event reporting records, i.e. AE, SAE.
- All record layouts are versioned and can be updated at any time during the trial.
- Integrated multi-layered subject privacy management.

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Licensing

Evado clinical has four tiers of service that can be purchased on a per trial basis.

- **Lite**: a simple easy to user service for small trials a maximum of 200 subjects and 10 sites
- **Standard**: a service for medium sized trials or registries to a maximum of 500 subjects and 25 sites
- **Enhanced**: a service for large sized trials or registries with greater than 500 subjects and 50 sites
- **Enterprise:** a service for customers that are running large numbers of trials and or registries

Trials and Studies

- Evado's online record design tools, provide a simple, yet powerful environment for designing Case Report Forms, Adverse Event Forms, Serious Adverse Event Forms, Protocol Deviation and Protocol Violation reports.
- All trial design components are fully versioned and released without affecting any of the existing trial data.
- Evado's trial scheduling module can manage 20 different data trial or study data collection schedules.
- Evado has central document publishing functions to manage the distribution of site documentation.
- Trial subject data collection schedules can be updated at any time to handle changes to trial's schedule, or when the subject returns to treatment.
- Multi-layered validation and management of mandatory data collection is integrated into the platform.
- Protocol exception reporting is fully integrated into the data collection workflow.
- Clinical Outcome Assessments are fully integrated into the data collection workflow.
- Electronic Informed Consent is fully integrated into the subject data collection workflows.
- Remote Informed Consent enables participants to consent to trials without having to attend the clinic.
- Patient recorded observations and clinical outcome assessments are fully integrated into the patient's clinical trial data collection schedules.
- Integrated virtual visit consultations using secure video conference.
- Customers can design and run their own monitoring or data management.
- Automated notification of new or updated CRF values, SAEs, and protocol exceptions.
- Evado has medical device specific study features to streamline medical device trial data collection.

Registries

- Evado's online record design tools, provide a simple, yet powerful environment for designing Case Report Forms and COA questionnaires.
- All trial design components are fully versioned and released without affecting any of the existing registry data.
- Evado's trial scheduling module can manage 20 different data trial or study data collection schedules.
- Evado has a central document publishing function to manage the distribution of site documentation.
- Multi-layered validation and management of mandatory data collection is integrated into the platform.
- Clinical Outcome Assessments are fully integrated into the data collection workflow.
- Patient recorded observations and clinical outcome assessments are fully integrated.
- Virtual visit consultations are fully integrated into Evado Clinical.
- Integrated virtual visit consultations using secure video conference.
- Customers can design and run their own monitoring or data management that saves time and money.
- Evado has medical device specific study features to streamline medical device trial data collection.

Evado Informed Consent

Evado Informed Consent module is a fully compliant paperless app that ensures that participants are fully informed before they decide whether to enrol in a trial or study that and can be run as a stand-alone service or integrated into a clinical trial or study.

- Simplifies the consent process and simplifies roll out.
- Device agnostic.
- Can be used in clinic or remotely.
- Lower costs
- Unique options with versatility to use with multiple systems including paper.
- Allows for remote clinical trial research.
- Supports the Australian informed consent requirements.

Evado Clinical Outcome Assessment

Evado's Clinical Outcome Assessment (eCOA) module enables clinicians and patients to directly enter observations and complete questionnaires for the studies and trials they are participating in.

- Easy to configure
- If fill integrated into the trial scheduling
- Easy creation and completion of COA records
- Form layouts can be used across multiple trials and registries
- Adaptable for different types of trials and registries
- COA records can be collected in clinic or remotely
- User friendly intuitive access
- Flexible data collection, on a desktop or mobile device
- Extensive reporting

Evado Virtual Visit

Evado Virtual Visit module is a fully compliant system that enables the site to conduct subject consultations virtually. The virtual visit has inbuilt auditing and meets current EU and USA regulatory and privacy requirements including GCP, GPDR and HIPPA.

- Enables face to face remote trial and study visits
- Virtual visits are fully integrated into the trial and subject scheduling.

More information

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