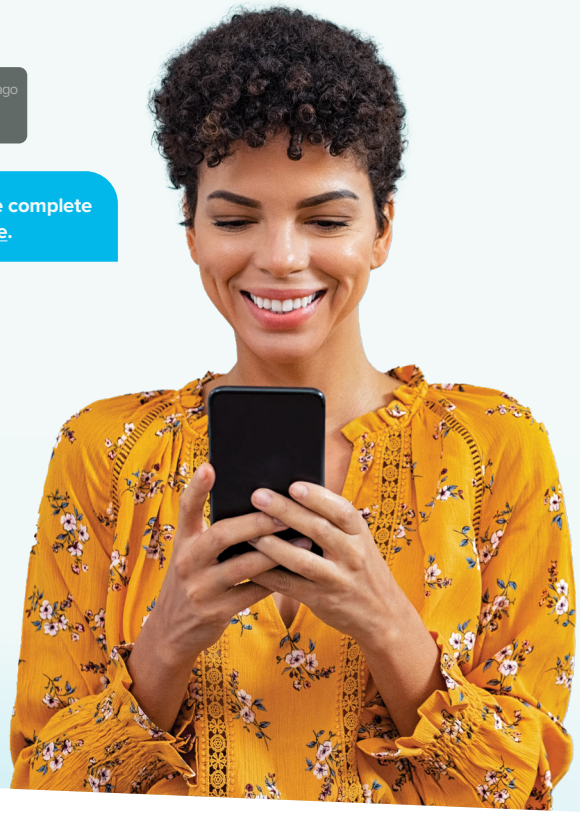
greenlight guru Clinical



Electronic Data Capture for MedTech Studies

The **leading** data collection toolbox purposefully built for MedTech.

Greenlight Guru Clinical is the only electronic data capture (EDC) system, purposefully built for MedTech. This SaaS offering allows users to collect and manage clinical data in pre and post-market clinical studies, including registries, cohorts, surveys, human factor testing, design validation, and more.

Our Customers Appreciate

Outstanding Support Team

★★★★★

(5 out of 5 Stars)

Pricing Subscription

User-Friendly Interface

Easy to Understand and Use Software

Why Greenlight Guru Clinical?

- Set up your first study in just 2 or 3 weeks vs. up to 12 weeks in other systems or even more using analogue methods.
- **Made for MedTech** = a modern, flexible platform that accommodates the MedTech workflow and operations and enables data collection throughout the whole life-cycle of a device.
- We meet the regulatory requirements of the FDA, EU, and most other countries.
- We ensure compliance out-of-the-bo with GCP, ISO 14155:2020, FDA cfr 21 part 11.

Get GG Clinical When?

- You need to run a clinical trial for CE marking, or PMA approval and you need to collect participants data in a compliant, secure way.
- You need to run post-market operations (vigilance, performance or PMCF)

greenlight guru Clinical

greenlight guru Clinical

eCRF

Avoid transcription errors, data entry costs, or data loss with our electronic Case Report Forms (eCRF). Design highly customized eCRFs in 3 easy steps.

ePRO and eCOA

Increase subject experience and fill out rates with Bring-Your-Own-Device (BYOD) electronic Patient Reported Outcomes (ePRO) via E-mail/SMS.

Surveys

Collect vigilance and safety data, clinical experience and feedback with our pre-validated surveys specifically designed for PMCF and PMPF.

Add Ons

Concomitant Medication | Register and track subject medication and link it to Adverse Event reports
Randomization | Randomize subjects using variated block size randomization, with or without stratification
AE/SAE Reporting | Comply with ISO14155 AE/SAE reporting requirements with ready-to-use GCP based templates
eConsent | Obtain electronic signatures on informed consents with remote signing options
Translations | Create forms and send questionnaires in 35+ languages
API access | Accelerates workflows, minimizes data inconsistency, increased cross system connectivity
SMS & notifications | Send questionnaires or subject events via SMS/email
Connected devices | Collect continuous data from wearables such as smart watches
White labeling | Add your company logo and name to SMS and emails you send to subjects
File Vault | Upload large files or images, up to a total storage space limit of 4TB

> “We’ve managed to cut weeks of work out of clinical tests simply by being more efficient in the way that we collect and manage data, with additional benefits of data being of a higher quality and error-free which will help us drive our products faster to market.”
>
> **Karen Elise Karlsmose Boel,**
> Head of Clinical Audiology and Usability, SIV - Oticon

Experience the Greenlight Guru Clinicial Difference.

Greenlight Guru Clinical offers a modern, flexible approach that accommodates the fact that clinical activities for MedTech are as different as the devices are many.

SEE THE DEMO →

greenlight guru Clinical