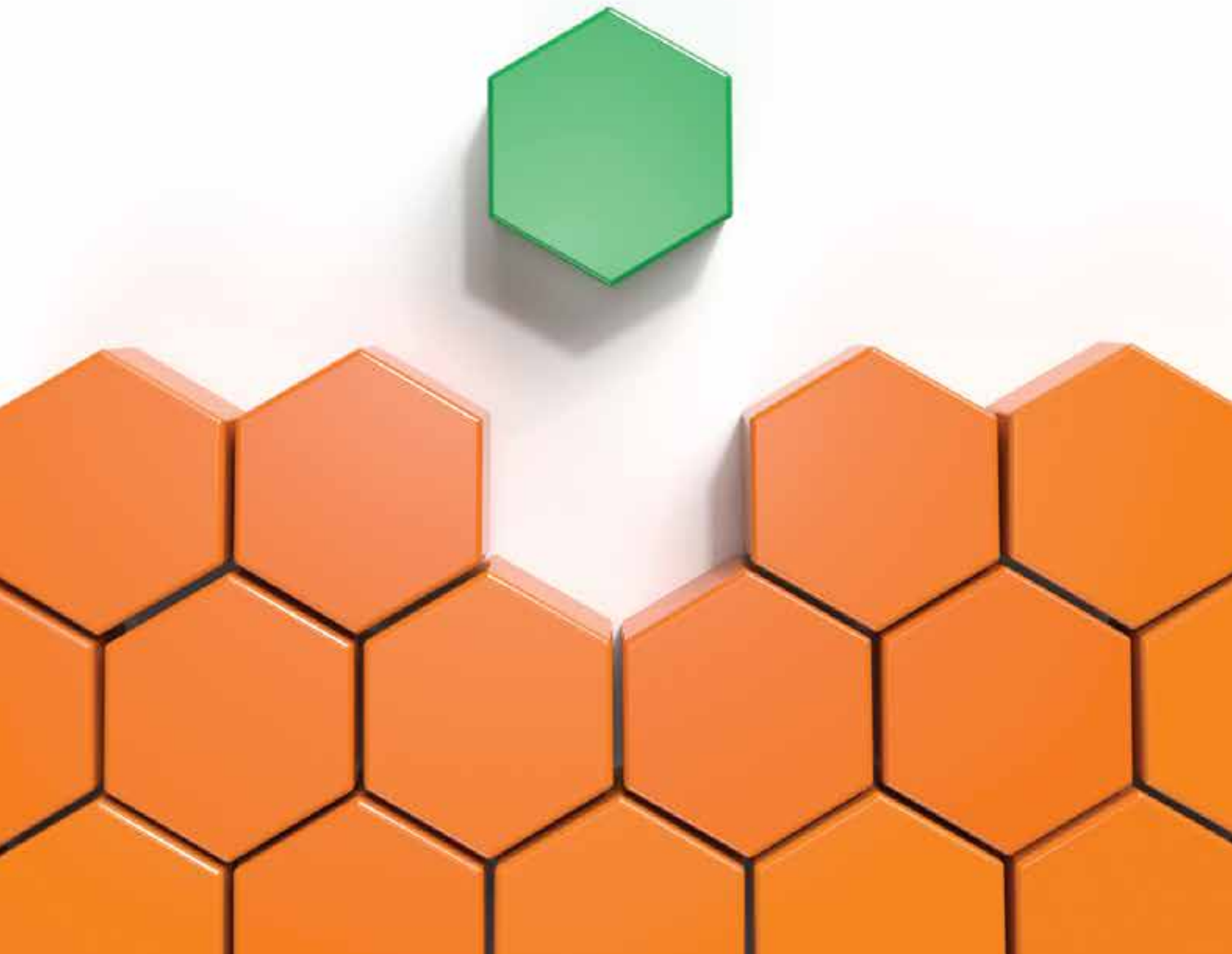
Viedoc™

eClinical solutions

Made to fit your trial



Simplifying clinical research since 2003

Viedoc is a comprehensive cloud-based platform built to improve the efficiency of conducting clinical trials. Created by industry professionals with decades of experience, Viedoc blends innovation with design to simplify your workday.

BENEFITS

Innovative design

Enjoy Viedoc's smart features and engaging design that allow you to focus on your research.

Quick start

Build your own study in no time or use our industry-standard templates.

Real-time reports

Have instant access to your data at any time with live reports and streamlined overviews.

Smooth workflows

Discover Viedoc's straightforward interface and experience more efficient workflows.

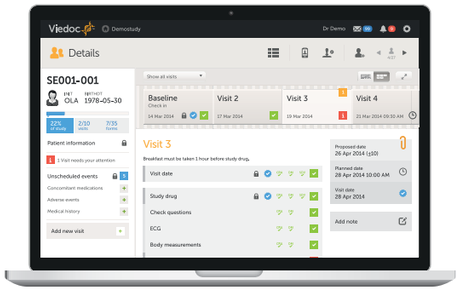
Fast data entry

Input data up to twice as fast compared to other EDCs with Viedoc's powerful system.

Flexible studies

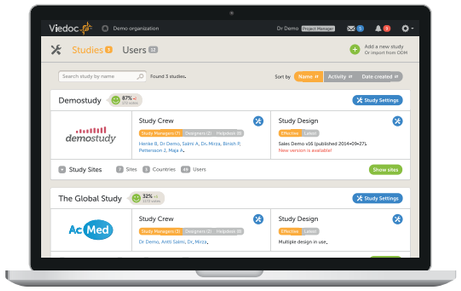
Make shifts and adjustments to your eCRFs as you go without needless delays.

Our eClinical suite offers a complete solution that is easy to use and simple to learn. Join Viedoc and spend time on what matters - **your research**.



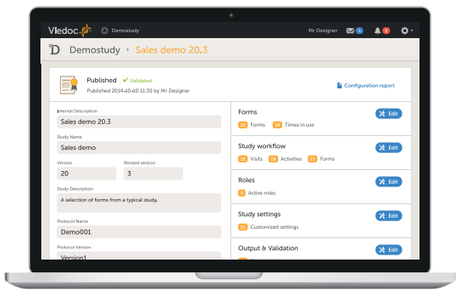
Viedoc Clinic

Data collection & monitoring made easy.



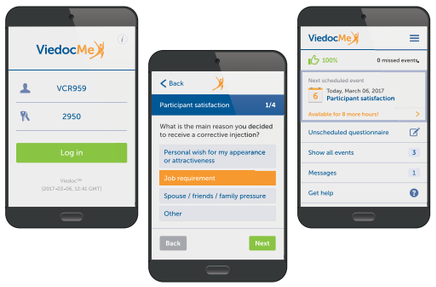
Viedoc Admin

Full control & study management at your fingertips.



Viedoc Designer

Professional study builds & libraries tailored to you.



ViedocMe

Patient input & reminders in one fully integrated app.

The most uncomplicated EDC ever experienced
- Novotech

Easy setup even for complex studies
- Aci Clinical

Flexible and feature rich
- ZIFO RnD Solutions

Excellent quality
- Ipsen

Really helps us save time
- Kuntuo

Extremely easy to use
- Neuromod

Shaping the future of EDC

Everyone is trying to keep up with technology. As a software company, you can either follow or lead. Our mission is to be at the forefront of eClinical innovation by developing new, smarter ways to help you succeed with your trial.

FEATURES OVERVIEW

Data Capture

- Web-based interface without client installation or client data storage
- Configurable role-based permission system

Data Management

- Realtime metrics & overviews
- Export to Excel, ASCII, SAS, PDF and CDISC ODM

Study Setup

- Drag & drop form builder
- Reuse of study building blocks & custom templates

Integrations

- Medical Coding Support with MedDRA, WHODrug, ATC & IDF
- API for integrations with other systems

Patient Reported Outcomes

- Native ePro integration including email & SMS reminders
- eCOA questionnaire library, support for VAS & photo upload

Data security

- Automatically backed up & securely protected data
- Encrypted communication & Two-Factor-Authentication

RTSM

- Randomization & advanced allocation
- Supply management & logistics with full inventory control

Support

- eLearning with online documentation & self-certification support
- Administrative interface for study maintenance & helpdesk tasks

Working for a healthier world

Viedoc was created to help life science companies bring therapies to market faster by developing more efficient EDC technology. In a world where minutes matter, we believe in eClinical software that makes a difference, one study at a time.

2 000+ STUDIES

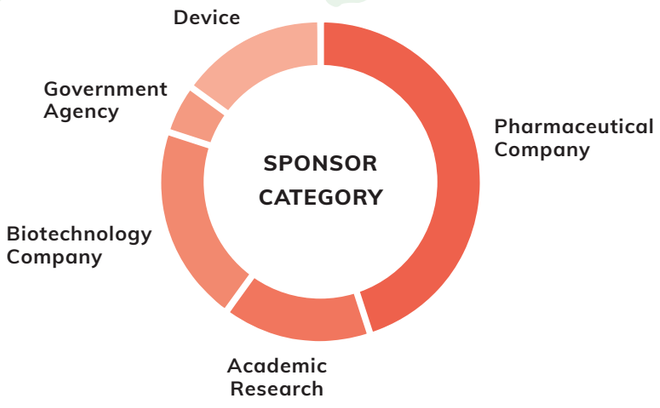
30 000+ SITES

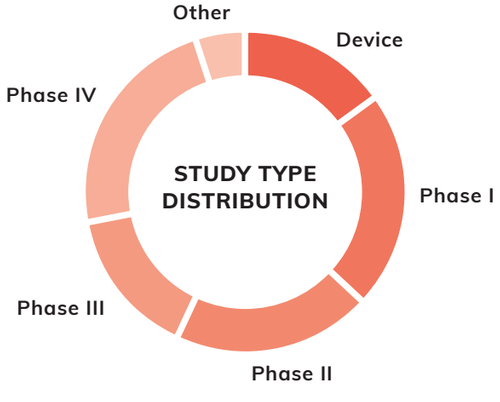
75+ COUNTRIES

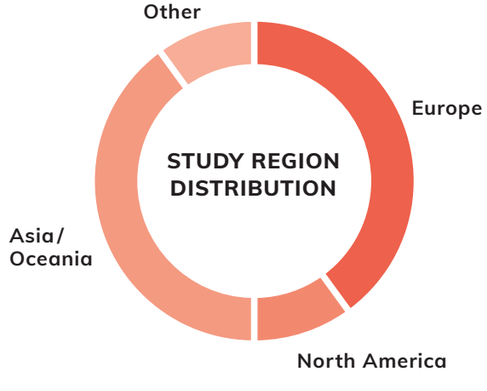
600 000+ SUBJECTS

300+ AUDITS

Bringing success to decades of trials







COMPLIANT & CERTIFIED

GDPR – EU law on data protection and privacy for all individuals within the European Union.

GAMP 5 – Framework for the risk-based approach to computer system validation based on the system's intended use and complexity.

21 CFR Part 11 – Establishes the FDA regulations on electronic records and electronic signatures (ERES).

CDSIC – Enables clinical research to work smarter by allowing data to speak the same language.

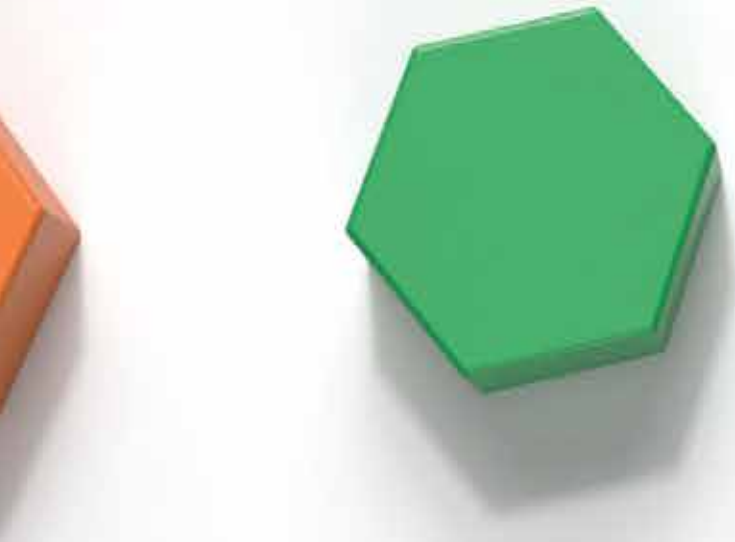
ICH GCP – Unified standard for the EU, JP and the US to facilitate the mutual acceptance of clinical data by the regulatory authorities in these jurisdictions.

APPI – Japan's Act on the Protection of Personal Information (APPI).

HIPAA – Health Insurance Portability and Accountability Act, sets the standard for protecting sensitive patient data.

PI-Specification – Information Security Technology – Personal Information Security Specification, China.

Enjoy your trial™



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