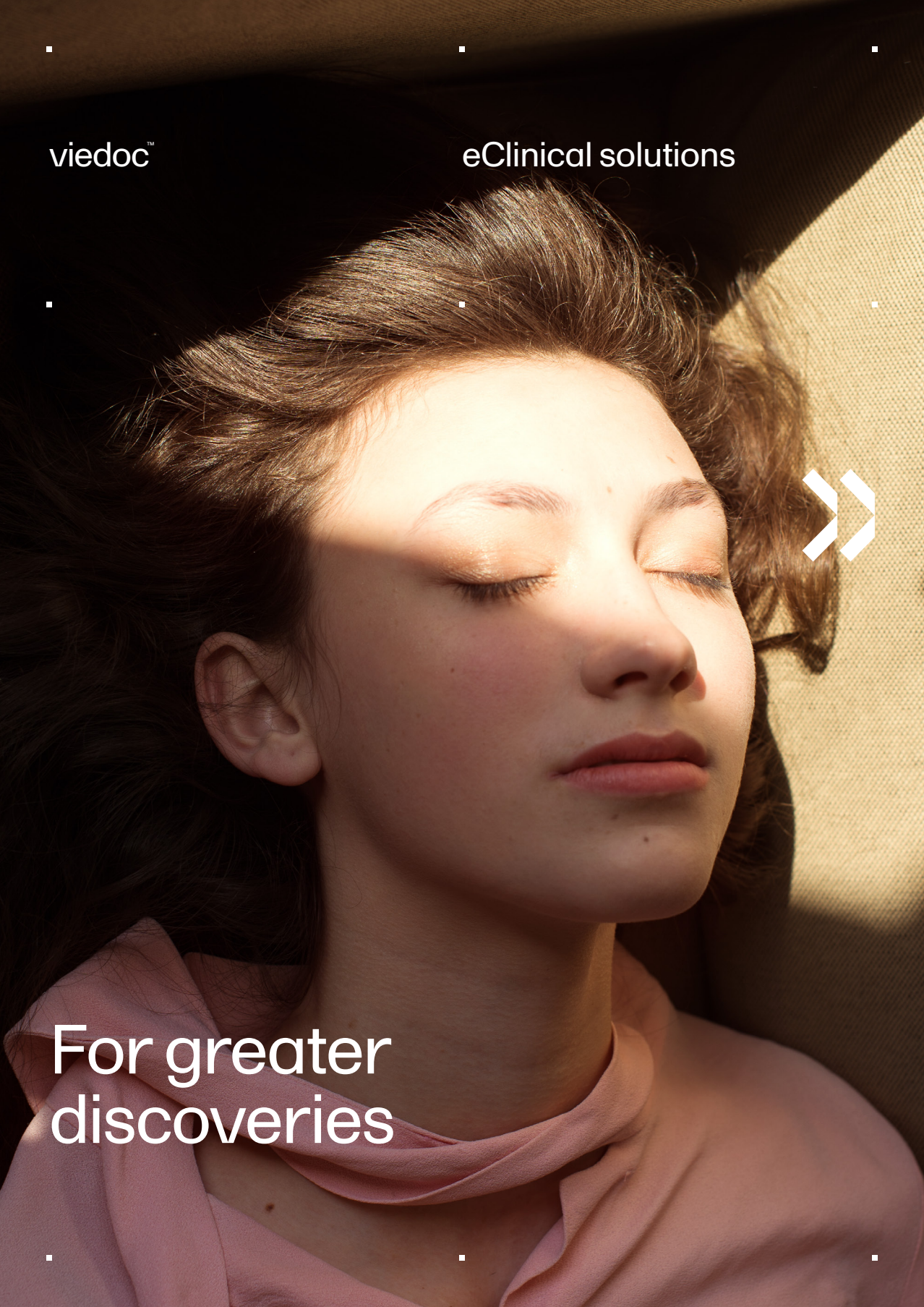
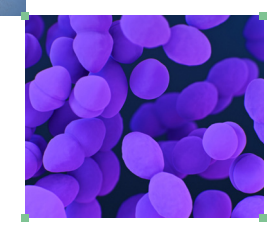
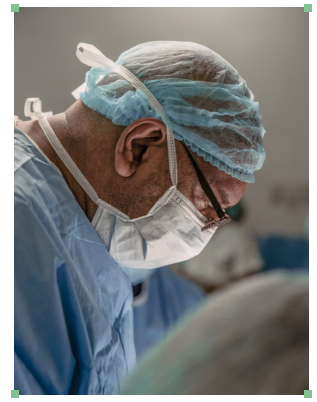

viedoc™
eClinical solutions
For greater discoveries

Accelerating clinical trials since 2003





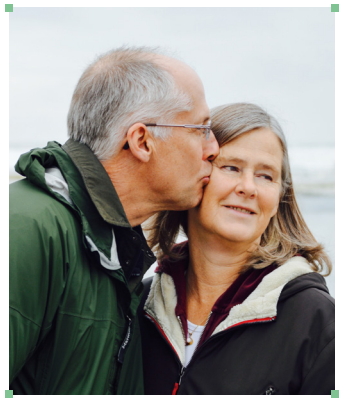
For two decades, we have perfected the craft of making clinical data more accessible so that greater discoveries can happen sooner.

Working for a healthier world

At Viedoc, we believe in life and science, in people, and in our collective power to change the world and build a healthier future. That unmatched driving force is what pushes us to innovate, accelerate and improve every aspect of modern clinical studies.

Since 2003, Viedoc has united scientists and clinical trial professionals in a shared mission of pushing life-changing research forward.

Our solution has been used to power thousands of studies, by collecting data from over a million patients and allowing it to flow smoothly across sites and countries. We take great pride in helping bridge the gap between patient and researcher – and, in the best of cases, between research and breakthrough.



Mirroring a rapidly changing world

Viedoc's eClinical solutions features the highest standard in data acquisition, management, and visualization. Whether for traditional multi-site studies or decentralized trials in a post-covid age, Viedoc is trusted by 9 of the top 10 large pharmaceutical companies for a reason.

The essentials

viedoc clinic™	For the investigator Manage all your trial data in one engaging solution
viedoc admin™	For the study manager Get your study started – and keep it running smoothly
viedoc designer™	For the study builder Create your own professional study – no advanced design or coding skills needed

The addons

viedoc me™	For the subject Reliable data collection, directly from the source
viedoc connect™	For the decentralized trial Fully integrated support for Televisits and eConsent
viedoc logistics™	For the supply manager Smooth, secure and seamless inventory tracking and randomization
viedoc tmf™	For the sponsor Powerful documentation management on investigator and sponsor level
viedoc reports™	For the data manager Tailorable reporting for quicker, deeper insights
viedoc pms™	For Japanese PMS studies Flexible data collection for the Japanese market

One ecosystem, endless possibilities

We offer the most engaging eClinical solution on the market. With Viedoc, you can design, manage and execute clinical studies in one scalable system. From there, pick and choose from a range of fully integrated addons – each specially developed to streamline a particular aspect of your trial.

Traditional, hybrid, or virtual. Viedoc adapts across studies, scales to each trial phase, and let's you collect data directly from the source.

Access what matters

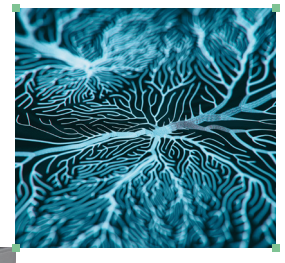
No matter how complex your clinical study may be, working with it should be smooth and simple. Viedoc combines an intelligent core with a streamlined interface and powerful tools, providing reliable access to user-relevant information – anytime, anywhere, from any device.

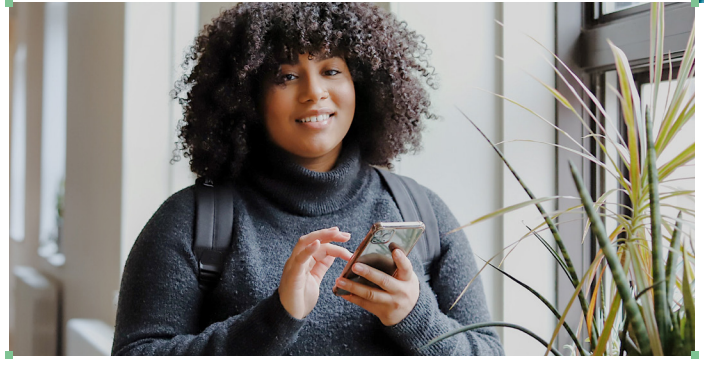
- Clean, straightforward interface
- Realtime metrics and at-a-glance dashboards
- Quick access to data – 24/7, 365 days a year

Accelerate every phase

Need for speed? Less time spent on setup, design, data collection and documentation means more time to focus on what's important: getting quality results.

- Web-based interface, requires no client installation
- Get your study build started in minutes, with ready-to-use, tailorable templates
- Collect data directly from subjects, by letting them submit information from their own devices





Adapt as you go

We see flexibility as an essential, not just a feature. Viedoc adapts across studies and scales to each trial phase. Traditional, hybrid, or virtual. And we've made it easy to adjust almost any aspect on the go, from design elements to user permissions – bringing complete control to your fingertips.

- Intuitive design tools and CDISC CDASH templates
- Make instant adjustments without interruptions or data loss – auto-backup and 24/7 protection
- Manage and configure all kinds of settings instantly – no tech department, designer, or helpdesk needed

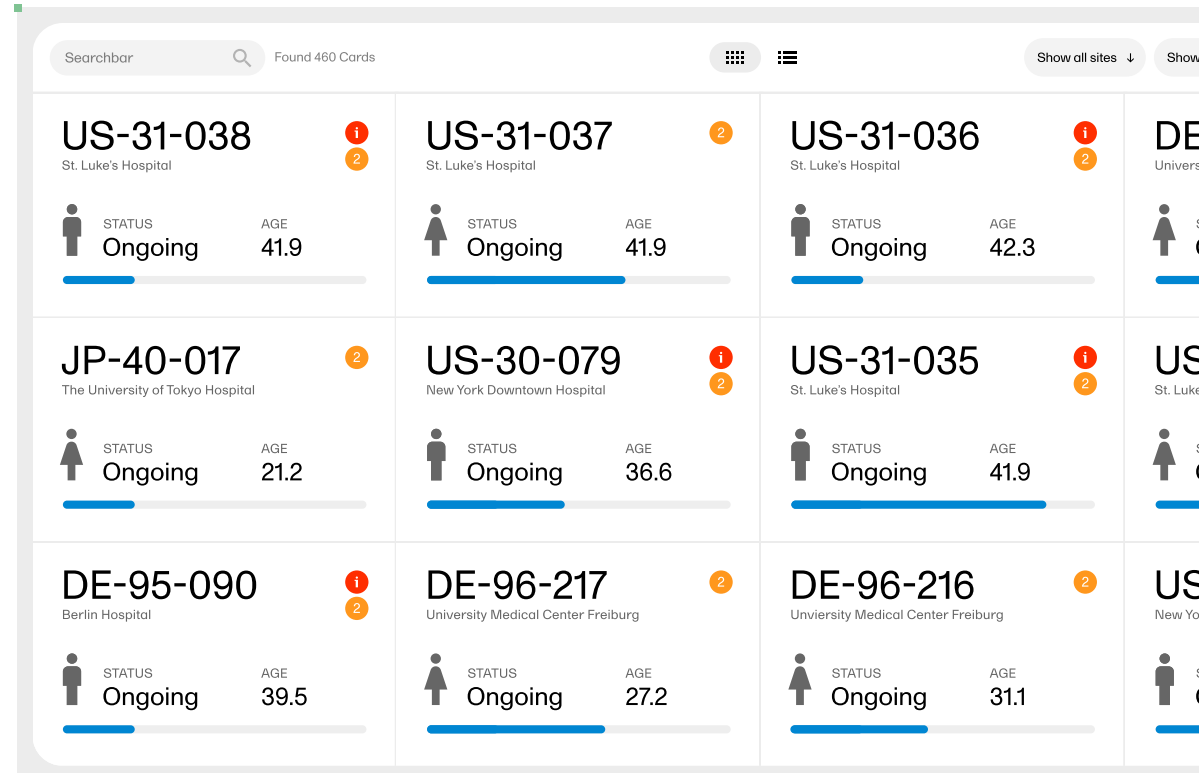
Trusted by 9 of the top 10 largest pharmaceutical companies.

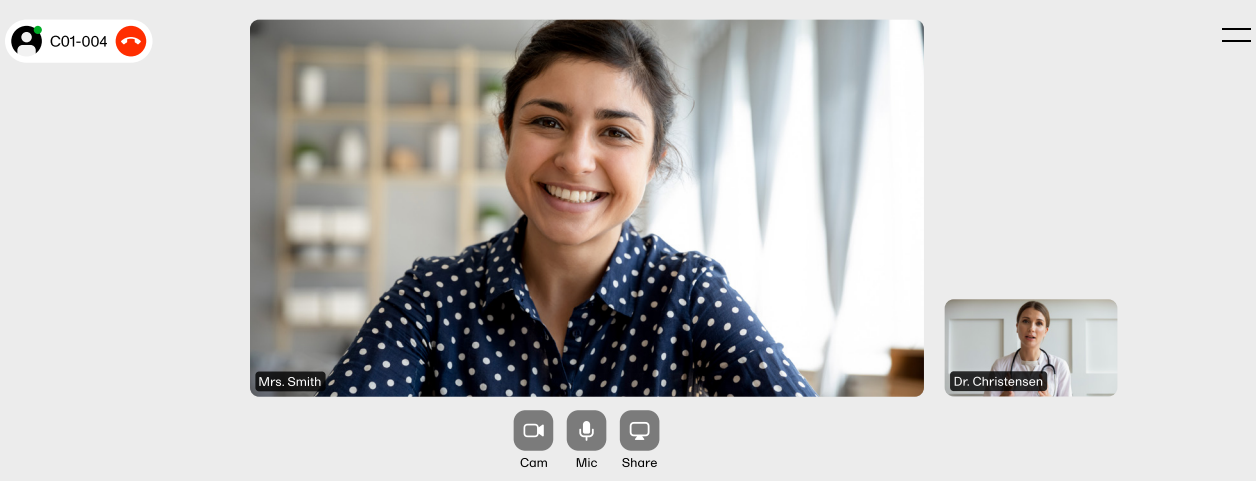
Viedoc at a glance

Viedoc exists to solve the data challenges of clinical trial professionals and improve how sites, CROs, and sponsors interact without ever compromising quality or safety. From being ISO27001 certified, ensuring the highest standards in information security, to having the fastest study setup times in the business, the benefits of choosing Viedoc are many.

One, unified solution

Move seamlessly between a complete set of applications and features covering all your needs – from setup to data delivery.





Remote

Full support for decentralized clinical trials – improving the subject experience.

Fast

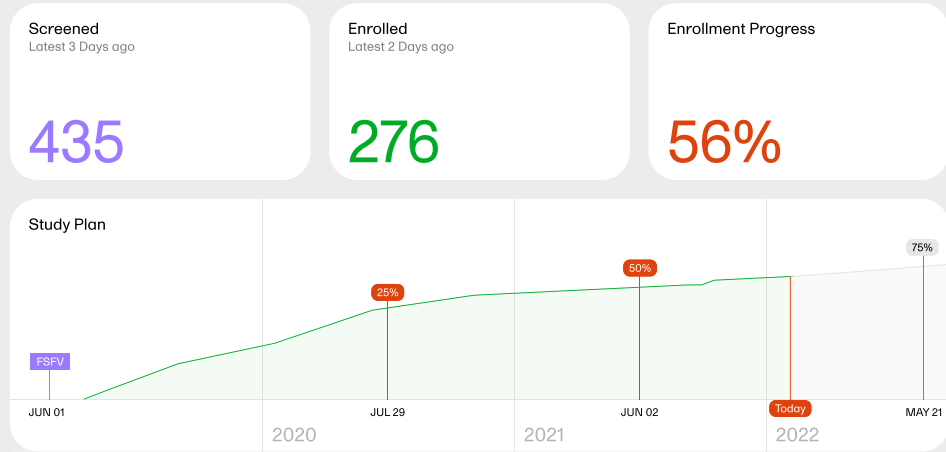
Accelerate setup times – Viedoc is 50% faster than other systems, from study setup to UAT-1.

Future-proof

New updates are released regularly, all backward-compatible. No additional system validation is required for new releases.

Flexible

Make mid-study changes at all levels with no system downtime in a self-service fashion.



Inspection-ready

Full documentation that meets inspection requirements, updated with every release.

Training for greater independence

Get started right away with our intuitive online guides, and beef up your skills with advanced study builder training.

No hidden costs

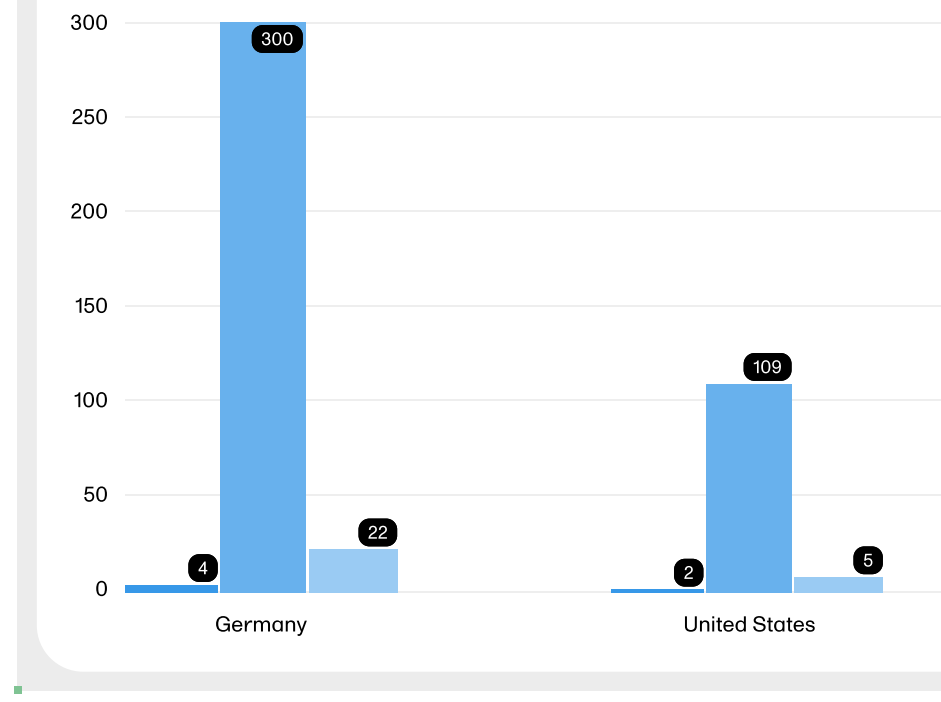
Pay as you go – no license fees until after the study starts, and no unexpected charges.

Global support

Access regional support and infra-structure, ensuring maximum performance wherever you choose to run your study.

Secure

All data is safeguarded using high-level security measures, including robust backup systems, advanced data encryption, and audit trails of all activity.



Users
100 000+

Sites
30 000+

Studies
4 000+

Countries
75+

Subjects
1 million+

Uptime
99.99%

Worldwide coverage

Viedoc's head office is in Uppsala, Sweden, with additional offices in North America, Europe, and Asia, servicing customers and studies in over 75 countries.

ISO 27001 certified

Viedoc has been granted the ISO/IEC 27001 information security management certification. This ensures our clients worldwide that our processes for implementing, maintaining and continually improving our information security management system (ISMS) follow the highest international standards.

100% compliant

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

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

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.

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

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

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

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

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

Want to know more about our solutions?
Ask for our product brochures, or download them at viedoc.com.

Viedoc designs engaging software for the life science industry. By accelerating clinical trials on all levels, our solutions support major pharmaceutical, biotech, and medical device companies, as well as renowned research institutions worldwide. Headquartered in Uppsala, Sweden, Viedoc also has offices in America, France, Japan, Vietnam, and China. Since our inception in 2003, over 1 million patients in more than 75 countries have participated in studies powered by Viedoc. Discover more at www.viedoc.com