



Clinical Trial Management Software Category

SUMMER 2026
Customer Success Report





Clinical Trial Management Software Category

Clinical Trial Management Software is a software solution used by research organizations, clinical research sites, and pharmaceutical companies to efficiently manage and streamline the various processes involved in conducting clinical trials.

Customer Success Report Ranking Methodology

The FeaturedCustomers Customer Success ranking is based on data from our customer reference platform, market presence, web presence, & social presence as well as additional data aggregated from online sources and media properties. Our ranking engine applies an algorithm to all data collected to calculate the final Customer Success Report rankings.

The overall Customer Success ranking is a weighted average based on 3 parts:

CONTENT SCORE

- ✓ Total # of vendor generated customer references (case studies, success stories, testimonials, and customer videos)
- ✓ Customer reference rating score
- ✓ Year-over-year change in amount of customer references on FeaturedCustomers platform
- ✓ Total # of profile views on FeaturedCustomers platform
- ✓ Total # of customer reference views on FeaturedCustomers platform

MARKET PRESENCE SCORE

- ✓ Social media followers including LinkedIn, Twitter, & Facebook
- ✓ Vendor momentum based on web traffic and search trends
- ✓ Organic SEO key term rankings
- ✓ Company presence including # of press mentions

COMPANY SCORE

- ✓ Total # of employees (based on social media and public resources)
- ✓ Year-over-year change in # of employees over past 12 months
- ✓ Glassdoor ranking
- ✓ Venture capital raised

Award Levels



MARKET LEADER

Vendor on FeaturedCustomers.com with substantial customer base & market share. Leaders have the highest ratio of customer success content, content quality score, and social media presence relative to company size.



TOP PERFORMER

Vendor on FeaturedCustomers.com with significant market presence and resources and enough customer reference content to validate their vision. Top Performer's products are highly rated by its customers but have not achieved the customer base and scale of a Market Leader.



RISING STAR

Vendor on FeaturedCustomers.com that does not have the market presence of Market Leaders or Top Performers, but understands where the market is going and has disruptive technology. Rising Stars have been around long enough to establish momentum and a minimum amount of customer reference content along with a growing social presence.

2026 Customer Success Awards

Check out this list of the highest rated Clinical Trial Management Software based on the FeaturedCustomers Customer Success Report.



* Companies listed in alphabetical order



2026

**CLINICAL TRIAL
MANAGEMENT
SOFTWARE**

MARKET LEADER





ABOUT ADVARRA



Advarra advances the way clinical research is conducted: bringing life sciences companies, CROs, research sites, investigators, and academia together at the intersection of safety, technology, and collaboration. With trusted IRB and IBC review solutions, innovative technologies, experienced consultants, and deep-seated connections across the industry, Advarra provides integrated solutions that safeguard trial participants, empower clinical sites, ensure compliance, and optimize research performance. Advarra is advancing clinical trials to make them safer, smarter, and faster.

96

TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“Advarra’s ICF writing services make our IRB submissions easier. The writers can work with whatever materials the sponsor provides, and unlike other vendors we’ve worked with, there’s less back-and-forth review time with the sponsor and the IRB. If Advarra writes the ICF, we don’t have to worry about it: we submit to the IRB and move on to the next task. Everything is just quicker with Advarra.”

REGULATORY MANAGER
CENEXEL CLINICAL RESEARCH

“Advarra has been able to significantly increase our clinical research budget funding, while drastically cutting our time dedicated to the budget process. We especially love how Advarra is able to cover the costs of their own fees. The Advarra team has been a fantastic partner for our clinical research program.”

CLINICAL RESEARCH DIRECTOR
CLINICAL TRIAL STARTUP COMPANY

“Advarra provides ACCRU with a thorough review process. This allows us to be more efficient with our study development and start-up times. The CIRBI platform is also easy to use and the customer service is excellent.”

BRETT PERCIVAL
COORDINATOR, REGULATORY AFFAIRS, MERIDIAN
CLINICAL RESEARCH

“Advarra eReg has been a huge time saver for our team and has improved regulatory compliance across our cancer center.”

ERIN NELSON
CTO REGULATORY MANAGER, THE OHIO STATE
UNIVERSITY COMPREHENSIVE CANCER CENTER

TRUSTED BY





ABOUT ARISGLOBAL



ArisGlobal is transforming the way today's most successful life sciences companies develop breakthroughs and bring new products to market. ArisGlobal end-to-end drug development technology platform, LifeSphere®, integrates ArisGlobal proprietary Nava® cognitive computing engine to automate all core functions of the drug development lifecycle. Designed with deep expertise and a long-term perspective that spans more than 30 years, LifeSphere® boosts efficiency, ensures compliance and reduces cost through multi-tenant Software-as-a-Service (SaaS) architecture. Headquartered in the United States, ArisGlobal has regional offices in Europe,...

22

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“ArisGlobal provided the best solution for compliance and to improve efficiency, and they have a clear roadmap that supports our future growth. We are very careful in choosing partners to ensure they live up to our high standards and fortunately ArisGlobal's safety database fits our needs very well. In fact, ArisGlobal was the only solution able to connect to the China FDA via gateway to meet their requirements.”

ANAND ANANTHAKRISHNAN
SENIOR DIRECTOR OF DRUG SAFETY AND
PHARMACOVIGILANCE, BLUEPRINT MEDICINES

“The collaborative relationship we have with ArisGlobal continues to be strong, and our joint plans for further enhancements reinforce our decision to choose them as our clinical technology partner.”

GLOBAL CRO PHARMACEUTICAL AND BIOTECH
INDUSTRY

“We needed a partner that would be attentive to our specific needs and concerns, and would ensure our voices were heard, and we found that with ArisGlobal.”

PAIGE QUEEN
SENIOR DIRECTOR OF CLINICAL OPERATIONS, TOTAL
CLINICAL TRIAL MANAGEMENT

“Traditional data entry and review processes using spreadsheet trackers and paper were no longer sufficient to ensure fast and high-quality submissions to the FDA.”

RACHEL RADWAN
DIRECTOR OF CLINICAL R&D, FUJIREBIO DIAGNOSTICS

TRUSTED BY



Syngene



ABOUT IQVIA



IQVIA (NYSE:IQV) is a leading global provider of advanced analytics, technology solutions, and clinical research services to the life sciences industry. IQVIA creates intelligent connections across all aspects of healthcare through its analytics, transformative technology, big data resources and extensive domain expertise. IQVIA Connected Intelligence™ delivers powerful insights with speed and agility — enabling customers to accelerate the clinical development and commercialization of innovative medical treatments that improve healthcare outcomes for patients. With approximately 79,000 employees, IQVIA conducts operations in more than 100 countries. IQVIA is a global...

269

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“IQVIA delivered invaluable support in preparing our organization for a critical new product launch. Their work helped clarify roles, responsibilities, and decision-making across our regional, global, and cross-functional teams. They enabled us to track all workstreams in a coordinated way, ensuring readiness for a successful launch. Their expertise in launch excellence frameworks was instrumental in pressure testing our plans and processes. We are grateful for their partnership in this high-stakes initiative.”

HEAD OF COMMERCIAL
LARGE BIOPHARMA COMPANY

“IQVIA’s AIML model project changed the organization and was in all my years probably the best thing I ever did. We were able to change how HCPs think about patients and realize a ~250% volume increase by uncovering all this additional opportunity.”

SR. DIRECTOR OF MARKETING
LARGE PHARMACEUTICAL BRAND

“[IQVIA NLP] provides data that would take tens or hundreds of times longer with tedious manual work. It enables downstream calculations to provide insight. Some work would not have been done or done comprehensively without [the platform].”

“Regulatory and HTA Considerations for Development of Real-World Data Derived External Controls.”

LEADING BIOPHARMACEUTICAL COMPANY

TRUSTED BY



Bayer HealthCare
Pharmaceuticals



EMORY
HEALTHCARE



ABOUT MEDIDATA SOLUTIONS



Medidata Solutions is the leading global provider of cloud-based solutions for clinical research in life sciences, transforming clinical development through its advanced applications and intelligent data analytics. The Medidata Clinical Cloud™ brings new levels of productivity and quality to the clinical testing of promising medical treatments, from study design and planning through execution, management and reporting. They are committed to advancing the competitive and scientific goals of their global customers, which include over 90% of the top 25 global pharmaceutical companies; innovative biotech, diagnostic and device firms; leading...

58

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“Medidata is a global brand that everyone knows, and everyone is happy to use. The seamless experience ... creates a much easier experience for us. Sites are entering data very quickly. They know the system and they know these modules back and forth. They don't have a lot of questions. And there's a lot of confidence in that.”

KARIS OASAN
DIRECTOR CLINICAL AFFAIRS, CONFORMAL

“The team of Professional Services that I work with is very intelligent and collaborative. We enjoy working with Medidata, it's a great team. They're always willing to come to the table and discuss our needs.”

HEIDI MCINTYRE
DIRECTOR OF CENTRALIZED DATA MONITORING,
MODERNA

“We needed to find a trusted partner to work with us, to grow with us AND deliver and provide innovative technology solutions for our clinical trials, we found that partner in Medidata.”

TESARO

“The Medidata products are very innovative and support our very complex oncology studies.”

VIJAY CHUNDRU
SENIOR DIRECTOR, EDC PROGRAMMING TEAM, JAZZ
PHARMACEUTICALS

TRUSTED BY

LUDWIG
CANCER
RESEARCH

moderna

Advanced Clinical

cognizant

DSP
CLINICAL



ABOUT VEEVA SYSTEMS



Veeva Systems is a leader in cloud-based software for the global life sciences industry. Committed to innovation, product excellence, and customer success, Veeva has more than 450+ customers, ranging from the world's largest pharmaceutical companies to emerging biotechs. Veeva is headquartered in the San Francisco Bay Area, with offices in Europe, Asia, and Latin America.

985

TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“By using Veeva Study Startup, we’ve reduced the overall timeline from site selection to site activation by 33%. And we shaved off six weeks from the time the protocol is approved to when the first subject is dosed.”

KURT BISCHOFF
ASSOCIATE DIRECTOR, BRISTOL MYERS SQUIBB

“Our clinical team was very excited because they handle the fallout when principal investigators or site users are not able to access a training or completing the signatures.”

DILLIRAJA PURUSHOTHAMAN
PRINCIPAL PROGRAMMER, FORTREA

“Archival is as important as trial operations because, when the study is over, all you have are these documents to show how the study worked.”

GERGANA KOUTSAROVA
TMF DIRECTOR, PROCESS QUALITY AND COMPLIANCE, ASTRAZENECA

“We’re partnering with Veeva to raise the standards across the industry and show what can be done within medical information.”

JENNIFER RIGGINS
PARTNERSHIP AND TECHNOLOGY STRATEGIST, PHACTMI

TRUSTED BY

abbvie

MERCK



Lilly

syngenta



2026

**CLINICAL TRIAL
MANAGEMENT
SOFTWARE**

TOP PERFORMER





ABOUT CASTOR EDC



Castor is a leading provider of clinical trial technology that automates the research process. Castor is helping Democratize Clinical Research and making DCTs scalable with EDC, eConsent, ePRO & eCOA + DCT integrations. These solutions enable researchers to easily capture and integrate data from any source and decentralize their trials.

50

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“Castor’s all-in-one platform was a game-changer. One coordinator runs our entire study, saving us an estimated \$100K in staffing while maintaining realtime data oversight. We’ve had nearcomplete questionnaire data, and the professional services team has been there every step of the way.”

MICHAEL LISS
ONCOBIOMIX

“Having an EDC system that can keep everything together is very important to successful research – especially when there are many variables like different languages, changes of shift, etc.”

ELIZABETH MILLEN
CLINICAL STUDIES PROJECT MANAGER, ADA HEALTH

“It’s easy and intuitive to build studies in Castor. The Castor Helpdesk with FAQs and the calculation tool were very helpful for study builds.”

JULIE ORELLINA
CLINICAL PROJECT MANAGER, OPTINA DIAGNOSTICS

“Castor allows us to be in control of our study and confidently run a study that follows good clinical practices, all at a fair price point.”

JAKE GILSON
ASSOCIATE DIRECTOR, MMI

TRUSTED BY





ABOUT CRIO



CRIO is a health tech company helping pharma, biotech, research sites, and academic research centers streamline regulatory workflows with one system that enables a single point of data capture. As the leader in eSource technology, CRIO enables remote monitoring and immediate data review with a powerfully integrated eSource/EDC solution. The system delivers a single point of data capture, eliminating the need to re-enter data into EDC.

63

TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“Using CRIO for our clinical trials has been truly transformational for our site. As a mobile site, we require flexibility in how we deliver trials and manage document storage, and CRIO has perfectly addressed these needs. The platform’s adaptability has streamlined our processes, making it easier to manage data, communicate with stakeholders, and stay compliant. We are grateful for the positive impact CRIO has had on our operations, and we look forward to continued success with this innovative tool. Thank you!”

MELISSA REIDY
SENIOR PROJECT MANAGER, HEART OF AUSTRALIA

“We chose CRIO to streamline the oversight and standardisation of operations across our community-based clinical site network. When looking for a site CTMS, we were drawn to CRIO’s intuitive user experience, which we felt would perfectly suit our community research teams, embedding quality by design across our organisation. The platform also stood out for the CRIO team’s implementation support, offering us a partnership approach as we grow our research portfolio.”

CLAIRE SEARS
HEAD OF BUSINESS OPTIMISATION, EMS HEALTHCARE

“ClinCloud was excited to collaborate on a study that integrated eSource data and EDC. With this capability, our site and staff can focus more time on recruiting and retaining patients – what we do best.”

JESSICA BRANNING
CEO, CLINCLOUD

“CRIO’s Reviewer interface is intuitive and easy to use, and with direct data capture, sponsors have a cost effective tool to take more clinical trial oversight in house.”

DAVID GOFFREDO
ASSOCIATE DIRECTOR CLINICAL PROJECTS, RXSIGHT

TRUSTED BY





ABOUT ECLINICAL SOLUTIONS



eClinical Solutions helps life sciences organizations around the world accelerate clinical development initiatives with expert biometrics services and the illuminate Clinical Data Cloud – the foundation of digital trials. Together, the illuminate platform and eClinical Biometrics Services give clients self-service access to all their data from one centralized location plus advanced analytics that help them make smarter, faster business decisions.

28

TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“We wanted to be digitally transformed and we really needed a centralized single version of the truth. Elluminate offered a cloud-based solution, rapid time to value with standard reports we got out of the box, and fertile ground to iterate over time based on our specific needs. Early on we needed a solution and elluminate fit that bill.”

RICH GLEESON
HEAD OF R&D TECHNOLOGY, CERVEL THERAPEUTICS

“The [eClinical Solutions] team’s knowledge of Karyopharm data standards and processes combined with their expertise in data collection platforms and clinical data management practices helped us meet our accelerated timelines for this important new research.”

KRISTAN GALLITANO
SVP DEVELOPMENT OPERATIONS, KARYOPHARM THERAPEUTICS

“Since implementing elluminate Data Central, there is no backlog in data, data review or queries in the studies that use the platform. Everything is addressed in a timely manner, both internally and from the eClinical team.”

JELLY DOMINIC MARATT
DIRECTOR, DATA MANAGEMENT, WAVE LIFE SCIENCES

“The elluminate platform helps to increase the confidence other Sarepta teams have in the data as we are all working from the same system, one centralized source of truth.”

ELIZABETH THOMPSON
ASSOCIATE DIRECTOR, CLINICAL DATA MANAGEMENT, SAREPTA THERAPEUTICS

TRUSTED BY





ABOUT ETHICAL GMBH



Ethical Cloud Software Services for Data Management in Clinical Trials. Working with Clinical Development departments they delivered so many applications including study tracking, data collection, queries management, coding, reconciliation, adjudication, protocol violations, laboratories, data review, narratives, supply management, documents sharing as many more. Ethical supports Endpoint Adjudication, Electronic Data Capture and eCOA / ePRO Patient Diaries with the user-friendly professional software solution designed to ease the Clinical Data Management...

21

TOTAL CUSTOMER REFERENCES

VIEW ALL REFERENCES

FEATURED TESTIMONIALS

“Using an online eAdjudication portal for our SERAPHIN study endpoints assessment improved the efficiency of the data collection and the quality of our processes, allowing a timely completion of the study. On-line management of Adjudication allowed rapid assessments of the study endpoints and greatly facilitated the work of the external clinical experts. The tool...”

LOÏC PERCHENET
DIRECTOR, GLOBAL POST-APPROVAL STUDIES GLOBAL MEDICAL AFFAIRS, ACTELION PHARMACEUTICALS LTD

“By using eAdjudication solution the Endpoint Adjudication Process can be managed 100% by the web based system, without any paperwork. All documentation and information will easily be managed within the system. The CEC members can log in to eAdjudication at any time and at any place. This is very efficient and we can have a great overview of the event status...”

ULRIKA BODÉN
TEAM LEADER OF THE CEC DEPARTMENT, UPPSALA CLINICAL RESEARCH CENTER

“With the eAdjudication® Endpoint Adjudication Process, documentation and information are managed 100% by the web-based system without any paperwork. The CEC members can log in to eAdjudication service at any time and at any place. This is very efficient, and we have a great overview of the event status, submission, and adjudication.”

PERNILLA HOLMGREN
COORDINATOR CLINICAL ENDPOINT COMMITTEE, UPPSALA CLINICAL RESEARCH CENTER

“Thank-you Ethical for all your hard work on the trial, the result is a great tool. I appreciated your training for the committee members, the demo was very engaging and provided an excellent kick-off for the adjudication committee.”

MULTINATIONAL SPONSOR COMPANY

TRUSTED BY





ABOUT FLORENCE HEALTHCARE



Florence advances clinical trials through software for managing document and data flow between research sites and sponsors. Florence eBinders is trusted by 7,200+ research teams to manage the eRegulatory and eSource for 1000+ active studies. Florence eTMF is the most flexible eTMF on the market designed for the Sponsor and CRO team wearing "multiple hats." Florence eHub is revolutionizing site to sponsor connectivity in a virtual site workspace for site inspection readiness oversight, remote monitoring, study startup, and quality control.

40

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“With StudyOrganizer, we can see which link matches up with which study, and get what we need faster. I don’t have to remember anything, it’s all right there for me. Before, it would easily take 15 minutes to find the link. But with StudyOrganizer, you just click.”

HILLARY ACUNA
STUDY COORDINATOR, ALSA RESEARCH

“I save at least an hour a day by using StudyOrganizer, which allows me more time to work on each of my trials; answering queries, recruiting, filing, and entering data.”

SUE SHIDEL
RESEARCH SUPERVISOR, ALLERGY & CLINICAL IMMUNOLOGY ASSOCIATES (ACIA)

“Everything is right there at your fingertips. It’s a definite time saver, and definitely more organized. It takes the confusion out of everything.”

CINDY SCHOELLER
REGULATORY SPECIALIST, ALSA RESEARCH

“The interface for eSignatures is so easy most of our PIs now take care of signing documents on their train ride in the morning.”

CENTRAL REGULATORY MANAGER
COMPREHENSIVE CANCER CENTER

TRUSTED BY





ABOUT GREENPHIRE PATIENT PAYMENTS



Greenphire is the leader in global clinical trial payment solutions. Greenphire’s best-in-class solutions optimize clinical trial performance by simplifying and streamlining payment processes from sponsors and CROs to sites and patients. Greenphire’s ClinCard and eClinicalGPS solutions easily handle any type of trial design and complexity, resulting in more accurate and compliant payments globally for both sites and patients. The choice of industry leaders worldwide, Greenphire provides better performance and better data, resulting in better trials

59

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“ClinCard has been a lifesaver and a timesaver. The hours it would take every week to process these payments cut into time I needed to spend on other projects, so I can’t say enough about the ClinCard program. I think it’s genius.”

CLINICAL TRIAL COORDINATOR
GLOBAL MEDICAL NETWORK

“Greenphire makes CHDI’s global payments faster, simpler, and more secure so that our study teams can focus their energy on our patients, our sites, and the clinical success of our studies.”

ROBI BLUMENSTEIN
PRESIDENT, CHDI FOUNDATION

“ClinCard is very easy to navigate, and user-friendly, allowing our team to jump right in.”

EMILY CAMERON
RESEARCH ADMINISTRATOR, MEDICAL AND SURGICAL SPECIALTIES, TUFTS MEDICAL CENTER

“ClinCard makes participant payments seamless and efficient.”

VICE PRESIDENT, PHASE I OPERATIONS
FULL-SERVICE CONTRACT RESEARCH ORGANIZATION

TRUSTED BY





ABOUT MEDABLE



Medable's mission is to get effective therapies to patients faster. They provide an end-to-end, global cloud platform with a flexible suite of tools that allows sponsors, patients, providers & CRO's to work together as a team in clinical trials. Their solutions enable more efficient clinical research, more effective healthcare delivery, and more accurate precision and predictive medicine. Their vision is to accelerate the path to human discovery and medical cures. They are passionate about driving innovation and empowering consumers. They are proactive, collaborative, self-motivated learners, committed, bold and tenacious. They are dedicated to making this world a...

28

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“In partnering with Medable, we look to combine Accenture’s extensive industry and professional services expertise with Medable’s proven and cutting-edge platform to deliver winning outcomes at the forefront of the digital age. Together, we will be able to help our life sciences partners accelerate clinical trials and improve the patient experience by scaling decentralized clinical trials across their enterprise.”

NICOLE PARAGGIO
STRATEGY MANAGING DIRECTOR, ACCENTURE

“Working with Medable and their eConsent platform allowed us to rapidly launch and successfully execute recruitment for a large, multi-facility clinical trial. Use of the platform was straightforward and their training demo modules made it easy for our organization to implement.”

GEOFFREY A. BLOCK,
ASSOCIATE CHIEF MEDICAL OFFICER & SVP OF CLINICAL RESEARCH AND MEDICAL AFFAIRS, U.S. RENAL CARE

“Accelerating the pace of clinical development will help patients get access to life-saving therapies more quickly. We’re excited to help power Medable’s work in accelerating clinical trials by making it easier to connect to the ecosystem of real-world data.”

TRAVIS MAY
CHIEF EXECUTIVE OFFICER, DATAVANT

“Our sites AND our teams are happy to report increased efficiency, reductions in ‘back and forth communication’, and less data entry.”

DIRECTOR OF CLINOPS
TOP 5 PHARMA

TRUSTED BY





ABOUT MEDRIO

medrio

Medrio has offered an integrated eClinical Software as a Service (SaaS) platform with a fully hosted Electronic Data Capture (EDC) system that drastically reduces study timelines and costs by putting study managers in control of their studies. By enabling studies to be built completely online without the need for any custom development, Medrio eClinical software allows studies to be ready in days instead of weeks. Medrio's hundreds of customers span across the world and include 12 of the top 20 global biopharmaceutical companies and 3 of the top 10 CROs.

81

TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“Medrio is just the easiest one to use. I’m not a programmer, but after just a couple of hours of training, I was able to start building. It’s just so easy to use, There is no need for migration, and everything flows well. I very rarely have problems, even when making mid-study changes, and I think it is user-friendly for the end users as well. Plus, the customer service is top-notch. Those are the best parts about Medrio, in my opinion.”

COURTNAY BUONOMO
ASSOCIATE DIRECTOR OF CLINICAL, ATLANTIC RESEARCH GROUP

“I’m a big fan [of Medrio]. I know how easy it is to work with, how easy it is to clean and get data out of our bid defense is introducing the platform to potential clients.”

MELANIE MARSHALL
SENIOR MANAGER OF DATA OPERATIONS, ONCOBAY CLINICAL

“With Medrio, we’ve got a much more effective workflow, allowing us to produce high-quality data without added headcount and focus on the research we are passionate about.”

ROSE RESEARCH CENTER

“Medrio does what they do very well. The way that Medrio listens to the users and implements the suggestions that they make is awesome — you don’t get that elsewhere.”

HERON THERAPEUTICS

TRUSTED BY





ABOUT OBVIOHEALTH



ObvioHealth is a global digital health organization providing real-world digital clinical trials that deliver better data mobile clinical study application allows for site-less trial design, allowing subjects to use their smartphones to participate in remote monitoring clinical trials from the comfort of their homes. For studies designed to include site visits, application allows for continuous data collection in between visits to the clinic.

28

TOTAL CUSTOMER REFERENCES

VIEW ALL REFERENCES

FEATURED TESTIMONIALS

“ObvioHealth has brought their deep clinical expertise and growing DCT capabilities to bear on our projects— always emphasizing a solution-based approach. Their learning agility, customer focus, and transparency make it easier to address nascent project challenges. The Obvio team is readily accessible, and they willingly share their knowledge. This has been a true win-win partnership.”

DENNIS JOSEPH
HEAD OF CLINICAL DEVELOPMENT, HALEON

“The new platform was a fantastic tool to keep track of enrollment. Communication with the team was seamless, and working with the team was very enjoyable. Overall, we were very satisfied with this collaboration and would be open to partnering on future studies.”

PHIL BLAZY
SENIOR R&D PROGRAM MANAGER, ENTRINSIC BIOSCIENCE

“When glitches arose, as would be expected in a novel situation such as this, the ObvioHealth team was available to work through the problems and correct them for subsequent patient entry. I'd feel very comfortable recommending ObvioHealth.”

TERRY PLASSE
MEDICAL DIRECTOR, REDHILL BIOPHARMA

“ObvioHealth helped us figure out how to keep patients engaged without giving them more migraines. Incorporating the live chat feature and notifications into the project has eased the connection for the participants.”

STEVE SCHAEFER
CHIEF EXECUTIVE OFFICER, MI-HELPER

TRUSTED BY





ABOUT ONESTUDYTEAM



OneStudyTeam, a member of the Reify Health portfolio, provides the cloud-based platform StudyTeam to accelerate the development of new and life-saving therapies. StudyTeam brings research site workflows online and enables sites, sponsors, and other key stakeholders to work together more effectively using common technology. The suite of StudyTeam solutions reduces site burden and helps sites pre-screen and enroll more patients, provides sponsors with end-to-end visibility into recruitment activity across all channels, and guides sites in conducting the trial for patients who have been enrolled. StudyTeam is trusted by the largest...

28

TOTAL CUSTOMER REFERENCES

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REFERENCES**

FEATURED TESTIMONIALS

“StudyTeam proved to be a valuable tool for the study. The team had real-time access to screening and enrollment activities whenever we needed it. We were able to utilize this information to make early and informed decisions which led to improvements to enrollment performance.”

RIDDHI CHANNA
SR. MANAGER, GLOBAL CLINICAL PROGRAM
MANAGEMENT, AMGEN

“OneStudyTeam is so involved, it never seemed like we were going to be stuck without support. The building of that relationship before we really jumped in all the way with eSource was very helpful and has been instrumental in the success of this for our site.”

ASHLEE MORRIS
LEAD CLINICAL RESEARCHER, SEACOAST KIDNEY &
HYPERTENSION SPECIALISTS

“[StudyTeam] gives us visibility into the comprehensive recruitment funnel that’s often very difficult for sponsors to see. It also gives us earlier visibility into how the recruitment platforms are performing.”

MIGNON KEATON
SENIOR CLINICAL PROJECT MANAGER, PROKIDNEY

“StudyTeam is a tool coordinators dream about. This is such a great system. It has everything we need in one easy-to-use place. We usually struggle to keep track of this information. Thank you so very much for bringing this system into my life.”

MERCK

TRUSTED BY





ABOUT STARLIMS



STARLIMS knows how to take the complexity out of managing quality initiatives. By helping their customers drive more revenue faster, achieve and maintain a manageable cost base, and reach their compliance objectives, they can focus on what really matters – getting products to market faster.

72

TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“Having STARLIMS as one of our LIMS offerings was really a nice fit. The STARLIMS system has a lot of capabilities that come with the system out of the box. However, the ability to customize those functionalities, that was one of the key pieces that attracted us to STARLIMS especially for more traditional analytical labs.”

CHRISTOPHER DEHEN
INFORMATION & DATA SCIENCE TECHNICAL MANAGER,
DUPONT

“The data collection and reporting efficiencies of STARLIMS help us meet our customer needs. From the customer perspective the STARLIMS system adds a level of professionalism to the lab. STARLIMS helps us deal with the pace of our business and allows us to be efficient.”

TOBY POLHAMUS
SENIOR DIRECTOR, MANUFACTURING, COTT

“STARLIMS has been instrumental for Bayer in simplifying compliance, offering meticulous audit trailing for every laboratory interaction within the LIMS.”

TASSILO STEFFL
SENIOR BUSINESS SOLUTION ARCHITECT, BAYER AG

“We love the STARLIMS system’s flexibility.”

LAB ENGINEER
SASOL

TRUSTED BY





ABOUT VIEDOC

viedoc™

Viedoc is design engaging software that modernizes clinical research so that necessary treatments can reach the people who need them faster. They accomplish this by combining technology and a creative design to streamline, automate and simplify conventional processes within clinical research. Simply put, Viedoc makes every aspect of a clinical study a bit smoother, resulting in greater discoveries that make a difference, and has the potential to improve lives.

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TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“The Viedoc system is very easy to navigate and develop, particularly for a team of naïve developers. The team at Viedoc were extremely supportive throughout our development process. Developing the application ourselves allowed us to implement the various innovative aspects of our trial and make changes in an efficient manner. Our team of clinicians finds the application extremely easy to use and the flexibility meets the requirements of our studies.”

SHONA D'ARCY
CHIEF SCIENTIFIC OFFICER, NEUROMOD DEVICES

“Viedoc and LINK Medical worked together on the proposal, with both parties contributing ideas to exactly how the eTMF should work and why. The developers were then able to take those ideas and turn them into specifications for the project, put it into production, and develop the eTMF.”

CLAUDIA WENNBERG
MARKETING AND BRAND MANAGER, LINK MEDICAL

“Viedoc allows us to offer a comprehensive service package to efficiently organize and manage all aspects of multi-center clinical trials in a manner that fulfills all GCP requirements and meets the needs of our customers.”

ANTJE HOPPENHEIT
CHIEF OPERATING OFFICER, PHARMTRACE

“Viedoc already provides the CSV files with the script to import it into SAS, so it's very easy for us to implement that.”

CORNEA VENTER
HEAD OF DATA SCIENCES, TECRO RESEARCH



TRUSTED BY





2026

**CLINICAL TRIAL
MANAGEMENT
SOFTWARE**

RISING STAR





ABOUT CLINIOPS



CliniOps provides an Enterprise Platform for the Life Science industry, enabling Hybrid and Decentralized Clinical Trials (DCT), Real World Evidence (RWE), Patient Quality of Life (QoL), Patient Engagement, powered by Integrated Telemedicine, with a vision to make drug trials accessible, inclusive, faster, and efficient.

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TOTAL CUSTOMER REFERENCES

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FEATURED TESTIMONIALS

“Deciding to proceed with our clinical study on prostate health at the height of the pandemic was very difficult and risky. Our partnership with CliniOps was significant in making it possible for the study team to safely enroll patients through eConsent and for data to be reliably and securely gathered and stored in a cloud-based repository. The study was completed and outcomes demonstrated Crila's effectiveness in promoting prostate health. In no small measure, do we attribute the successful completion of our studies to the innovative use of CliniOps technology.”

SUE MCKINNEY
EXECUTIVE DIRECTOR, CRILA HEALTH

“CliniOps customized their system to organize data collection for our comprehensive community health screenings. Our events collect demographic, physical measurement, and point-of-care laboratory data from blood and urine testing. CliniOps tablet application was very intuitive and a quick 15 min onsite training was enough for our volunteers to start using the system seamlessly. The team was able to record about 200 patient data, in less than 4 hours, without issues. Excellent customer service.”

JAMES IVIE
DIRECTOR, NATIONAL KIDNEY FOUNDATION OF ARIZONA

“CliniOps is innovative, easy to use, perfectly positioned and has the potential to transform how we recruit and consent subjects. I believe that it is a real innovative improvement in clinical research.”

JOHN MENDELSON
CALIFORNIA PACIFIC MEDICAL CENTER RESEARCH INSTITUTE

“CliniOps App is a really cool application to make source documentation in any hospital, for research or clinical trials, really easy and user friendly. We are using them on a few trials at NH and find it quite interesting.”

ALBEN SIGAMANI
PRINCIPAL INVESTIGATOR, NARAYANA HEALTH

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ABOUT CRUCIAL DATA SOLUTIONS



Founded in 2010, Crucial Data Solutions (CDS) provides the most innovative and affordable data collection and clinical trial management technology on the market today. TrialKit, their cloud-based platform available via a web interface and downloadable native mobile app, enables end-to-end clinical trial management for pharmaceutical, biotechnology, and medical device companies of all sizes. Design and deploy regulatory compliant studies in days instead of weeks using their intuitive forms designer that requires no programming. Over 10,000 global users have leveraged the flexibility of TrialKit to deploy over 1,100 studies across all phases of clinical...

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TOTAL CUSTOMER REFERENCES

**VIEW ALL
REFERENCES**

FEATURED TESTIMONIALS

“The customer support provided by the TrialKit team was exceptional. They are responsive, knowledgeable, and always ready to assist with any questions or concerns. Knowing I have a dedicated support team backing me up was reassuring. Overall, TrialKit has simplified and automated my trial management process, saving me time and effort while delivering excellent results. It's a flexible tool for a start-up or larger trial, and I...”

HEAD OF CLINICAL OPERATIONS
CLOUDCATH

“I have built databases using SAS and Oracle Clinical, and have used many other systems. TrialKit is by far my favorite. The Customer Support is better than anything I've seen, literally, they are always just a phone call away and go above and beyond to make sure their clients are happy. Crucial Data Solutions is always seeking to improve their products and are always ready...”

LAURIE JOHNSON
CLINICAL DATA MANAGER, ICON GOVERNMENT AND
PUBLIC HEALTH SOLUTIONS (GPHS)

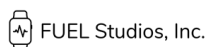
“TrialKit is an affordable, flexible, easy to use, integrated system. It allows us to reduce study build times and helps us to deploy study databases in a more timely manner for our clients. The support staff is extremely responsive and knowledgeable, and they are always seeking ways to improve their products. We have a very collaborative working relationship with the team at CDS.”

VP OF GROWTH STRATEGY
LEADING MID-SIZED CRO COMPANY

“TrialKit is an affordable, flexible, easy to use, integrated system. It allows us to reduce study build times and helps us to deploy study databases in a more timely manner for our clients. The support staff is extremely responsive and knowledgeable, and they are always seeking ways to improve their products. We have a very collaborative working relationship with the team at CDS.”

SVP RESOURCING OPERATIONS
CATALYSTCR

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ABOUT ENNOV



Ennov Group is an Independent Software Vendor with Electronic Document Management (EDMS) & Business Process Management (BPM) for Life Science Industries. Ennov Solutions, Inc. is a software publisher specializing in Quality Management Systems, Regulatory Compliance Solutions and Clinical Trial Management Software with more than 450 clients worldwide and 15 years experience in Pharmaceutical, Cosmetic and Healthcare Industries.

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TOTAL CUSTOMER REFERENCES

[VIEW ALL REFERENCES](#)

FEATURED TESTIMONIALS

“The Ennov project was very successful and a great experience. The aggressive timelines made the project very challenging. What kept us going to make the finish line was the great sense of teamwork between the two organizations. I think we learned a lot from each other and we will continue to learn from each other.”

SENIOR IT MANAGER
US-BASED MULTINATIONAL HEALTHCARE COMPANY

“It’s really nice to work with a software publisher that shows a great deal of innovative spirit, has great ambitions and is proactive. Above all, I really appreciate Ennov placing particular emphasis on the listening experience of the customer.”

CYRILLE JEUNE
PROJECT LEADER, AGUETTANT

“What sets Ennov apart is the unified database. You could integrate your eTMF with your clinical database and submission documents because the data is all in one place.”

JAKOB JUUL RASMUSSEN
FOUNDER AND MANAGING DIRECTOR, PHARMA IT

“The Ennov project team adapted to our expectations and rolled up their sleeves to meet tight deadlines.”

HORIBA

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HORIBA BACHEM