



Zelta: Making the future of clinical trials simple

Control at every stage. **Confidence** in every outcome.

Zelta is an experienced, trusted, market-leading cloud-based SaaS platform for modern clinical trials.

Zelta puts you in full command of every aspect of your clinical trials and research — from designing workflows and forecasting study costs, to building diaries that your participants can easily and intuitively fill out. Our vision is to provide dynamic solutions that drive bold research.



Choose a platform designed to scale and accelerate your trials

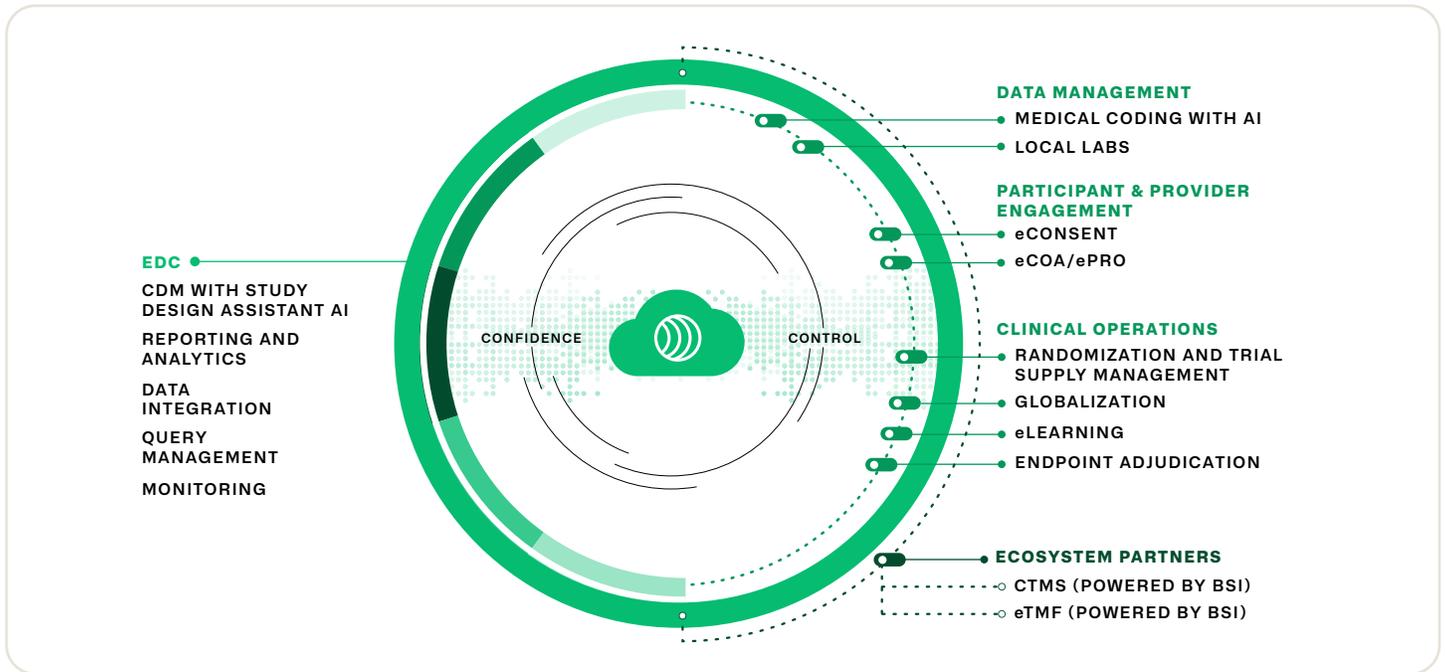
Zelta, a unified cloud-based clinical data management and acquisition platform with customizable modules, can be tailored to meet the unique needs of your clinical trials and accelerate outcomes.

What you get

Confidence in every outcome with a robust EDC

What you choose

Control at every stage with fully integrated modules



Zelta is the platform of choice for clinical trials



Flexibility & scalability: Manage clinical studies and research across all phases, therapeutic areas, levels of complexity, and geographic locations.

- For over 10 years, Zelta has been used in more than **4,300 clinical trials** from startup to submission, across all phases of clinical research, including **over 500 phase III trials** and **23 therapeutic areas**.
- Execute studies on a flexible **cloud-based EDC/CDMS platform** designed to manage complex study requirements with sophisticated trial design capabilities.
- Select from customizable modules, built on a **unified code base**, that allow you to leverage the functionality you need for each trial. Enable real-time access across the platform without needing to integrate disparate systems.



Ease of use: Implement and execute research, manage participants, assign tasks, and report results to stakeholders from a single, user-friendly interface.

- Reduce EDC build and validation time on even the most complex oncology builds through our **automated study processes** that streamline repetitive tasks, simplify cycle creation, and make it easier to manage recurring study elements.
- Use **supervised machine learning** to streamline the manual process of annotating CRFs with CDASH to reduce manual work and ensure accuracy.
- Leverage **real-time edit checks** to alert users to inadvertent data entry, reducing the cost of generalized query-based data cleaning.
- Get the right data for the right person using our **built-in intuitive report designer** that lets you build, define, and filter data and reports for specific roles and individuals.

Built with all users in mind, Zelta modules are fully integrated and share one code base with the rest of the unified platform – streamlining clinical trial processes and maximizing patient, caregiver, and provider engagement to accelerate clinical trial outcomes.



EDC

Design, validate, and launch studies and apply amendments without database migration



Data integration

Build and automate data connectors with minimal coding



RTSM

Streamline processes so you can manage with ease



Globalization

Ensure site data collection and ePROs are accessible, accurate, and compliant across multiple languages and regions



eLearning

Manage study training electronically to ensure regulatory compliance



Medical coding with AI

Dramatically streamline the coding process, enhance accuracy, and ensure compliance with regulatory standards



Local labs

Ensure data accuracy and integrity by standardizing lab data across sites when frequent sampling and fast results turnaround are essential



eConsent

Deliver quick and easy remote consent with no integration



eCOA/ePRO

Engage directly with participants and caregivers via in-app assessments and real-time analysis



CTMS (Powered by BSI)

Deliver comprehensive, easy-to-use functionality for all aspects of your clinical trials



eTMF (Powered by BSI)

Everything you need to capture and maintain regulatory binders and documents



Seamless access: Use a single sign-on from anywhere in the world to access a unified cloud-based solution built on one code base.

- Mix third-party solutions with Zelta and our ecosystem partners with **plug and play APIs** to create a truly configurable platform.
- Leverage Zelta study metadata, such as forms and event schedules, in **third-party eSource integrations**.
- Eliminate manual data entry by pushing **eSource/EHR data** to electronic case report forms.
- Enjoy a complete, seamless experience with **integrated CTMS (powered by BSI), eTMF (powered by BSI), and CDMS technology** – or easily integrate the solutions of your choice.



Happy users: Industry-leading CSAT scores of **>95%** prove that sponsors, CROs, sites, and participants prefer Zelta over other platforms.

- Trusted by more than **200 biopharma, medical device, CROs and AROs**.
- Call upon certified, experienced designers for support, **24/7/365**.
- Supporting **over 100 countries**, speaking over 70 languages and dialects.
- Zelta's **consulting, enablement, and extension services** are all tailored to users' needs and help them overcome any challenges they face in their clinical trials.

Awards and recognition

Zelta is ranked as Leader in 2024 Everest Group's EDC Matrix® Assessment.

[Read more about our ranking.](#)



About Zelta

Zelta by Merative is a clinical trials solution business that includes both a clinical data management and acquisition platform and consulting, enablement, and extension services. Zelta's unified cloud-hosted platform supports all phases and complexities of research, including more than 500 phase III trials.

Learn more at merative.com/clinical-development

About Merative

Merative provides data, analytics, and software for healthcare and government social services. With focused innovation and deep expertise, Merative works with providers, employers, health plans, governments, and life sciences companies to drive real progress. Merative helps clients orient information and insights around the people they serve to improve decision-making and performance.

Learn more at merative.com



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