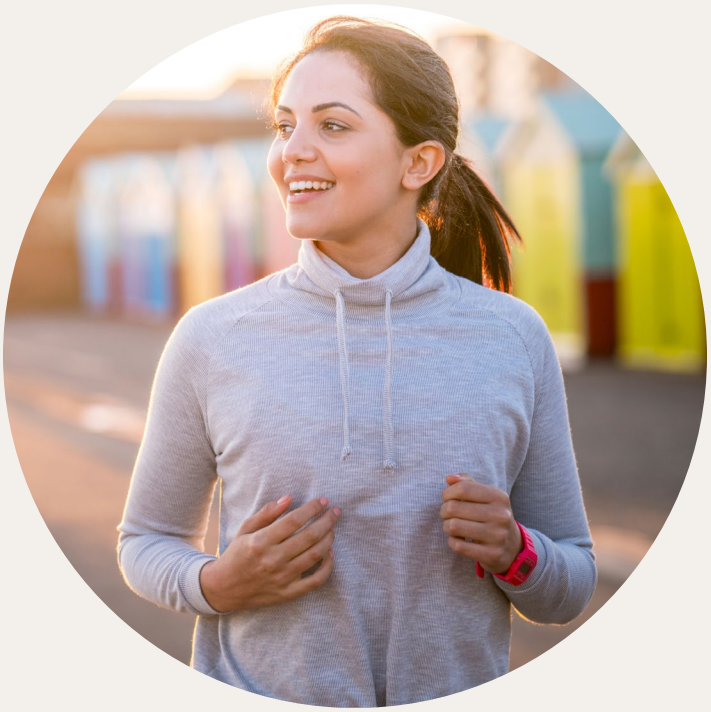
Zelta



FOR MEDICAL DEVICE COMPANIES

Zelta: a clinical trials platform

Control in every stage. Confidence in every outcome.

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

With the Zelta clinical trials platform by Merative, you are in full command of every aspect of your medical device trials and research — from designing workflows and forecasting costs, to building diaries for your participants.

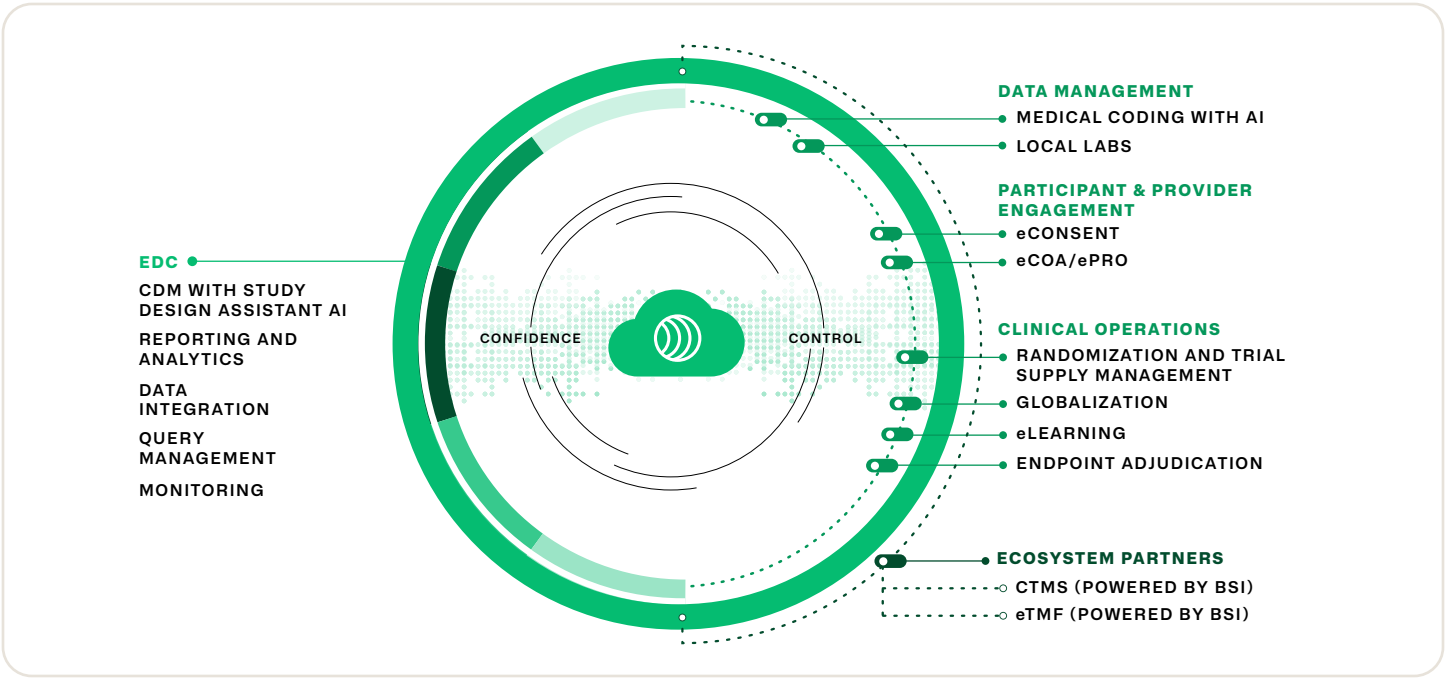
merative

Choose a platform designed to scale and accelerate your medical device trials

Zelta, a unified cloud-based clinical data management and acquisition platform with customizable modules, can be tailored to the 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 medical device trials

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

Scalable

- Control research of all types regardless of phase, therapeutic area or geographic location
- Host and scale thousands of trials around the world
- Create, standardize and scale processes to optimize cross-study control and reporting
- Help maximize international sites and patient engagement, supporting 75+ languages and dialects

Intuitive

- Make it easier to implement and execute research, manage participant compliance, perform routine tasks and report results to stakeholders through a single, user-friendly interface
- Optimize for sites and users
- Design trials with zero programming knowledge
- Take direct control of study go-lives, protocol amendments, study design changes and study closeouts

Unified

- Access modules and reports through a unified platform from anywhere in the world with single sign-on and one code base
- Remain current and ensure all of your trials and users are on the latest version of code with our single-instance technology
- Streamline clinical trial processes and help maximize patient, caregiver and provider engagement with clinical operations and patient and provider modules

Why Zelta?

Services
Whether you prefer self-service or full-service solutions, Zelta offers technology and deep medical device industry knowledge to help you overcome common challenges in clinical trials, whether it would be MDR, 510(k), or other medical device needs.

Data security
Our unified platform is hosted on a secure and flexible HIPAA-enabled cloud.

Flexibility
From small local studies to complex trials on a global scale, Zelta offers flexible functionality and pricing plans that can be customized based on your needs.

Platform support
Our team of certified, experienced designers are here to support you 24/7/365.

Trusted global partner
Experience supporting all phases of clinical research for 15 years with study sites in more than 109 countries and across 23 therapeutic areas.

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

500+

Medical device trials have been executed with Zelta

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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Take the first step to boost the efficiency of your clinical trials. Speak with your sales representative or read more about Zelta at

merative.net/contact-zelta