



an inVentiv health company

www.i3global.com

# Superior clinical data solutions

i3 provides the pharmaceutical, device, and biotechnology industries with comprehensive, integrated data service solutions that include data capture, summary, analysis, and reporting for clinical trials across all phases of research. We demonstrate leadership and innovation by leveraging world-class process design and superior quality to maximize efficiencies and exceed customer expectations.

## Strategic global presence

i3 brings together the best people, processes, and technology on a global scale to deliver high-quality data, accelerating the clinical trial process while conserving your resources. Our dominant presence in many emerging markets allows us to operate in lower-cost locations without making any sacrifice in results—as proven by our large and growing data management and statistical programming group in India—and provide effective, economical solutions to your challenges anytime, anywhere.

## Industry-leading experience

Our approach to data services is also shaped by our industry-leading level of experience, which spans not only international boundaries but studies from Phase I to Phase IV. This experience enables us to precisely pair technology and proven processes with the needs of your study, and offer you expert assistance in streamlining the route to more effective answers.

i3 data services include:

- **Comprehensive data management** that spans eCRF/CRF design, database design and validation programming, data capture and validation, focused data review and reconciliation, data analysis and integration, and database lock and freeze. Our combination of technology expertise and data experience enables us to customize solutions, anticipate data issues and trends, and seamlessly support every aspect of a project.
- **EDC and integrated technologies** that allow us to select the most effective application for each unique trial. Our use of such tools as Oracle® Remote Data Capture (RDC) Onsite,

and KnowledgePort®, our secure web portal, are just the beginning of our fully integrated technology platform.

- **Biometric services** including statistical analysis and consulting, study design, protocol development, randomization, data safety and monitoring support, and regulatory submissions report and representation. Our statistical programming services include the creation of SDTM data sets, NDA-ready data listings, summary tables, statistical figures, and integrated safety and efficacy summaries. Our experienced adaptive design consulting group has the expertise to evaluate trial results and recommend midcourse adjustments to help trials succeed on a faster and more cost-effective basis.
- **Medical and scientific writing** including clinical study reports, brochures, summary documents, and scientific presentations and manuscripts. Our access to technical and scientific authorities throughout the i3 family with extensive medical, statistical, and data management expertise allows us to fulfill whatever needs emerge for manuscript development.

i3 has more than 17 years' experience providing top biopharmaceutical companies with high-quality data services. This track record—combined with best-in-class technology, a customer-focused approach, and continuous process and quality improvements—allows us to fulfill your most demanding needs.

Find out how i3's expert clinical data solutions can support your product.

## Contact us

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[www.i3global.com](http://www.i3global.com)