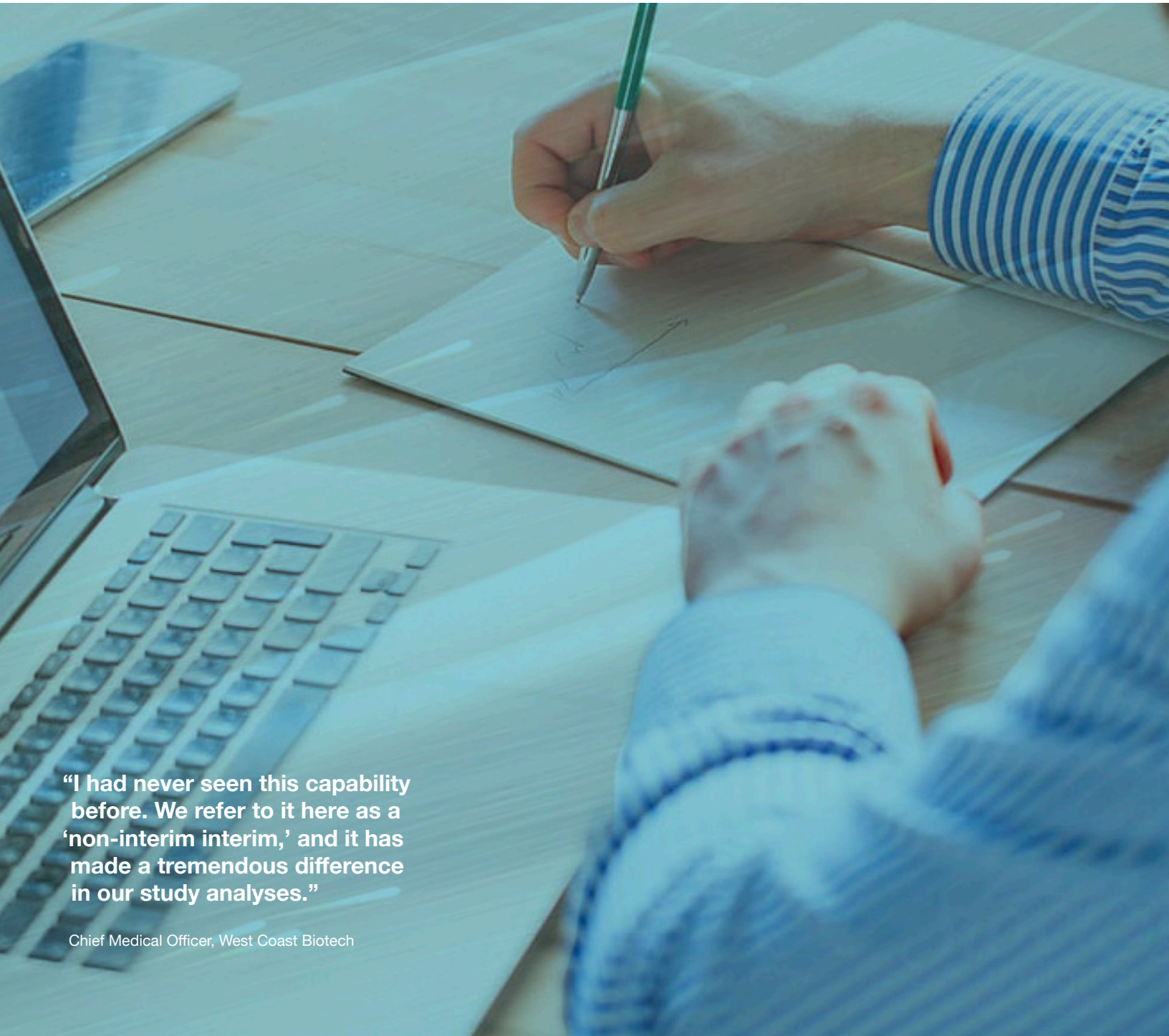
Prevail Integrated Clinical & Operational Analytics

PREVAIL INFOWORKS | RISE ABOVE

Prevail's innovative analytics provide timely, data-driven insights into live studies, regardless of protocol, data, and system complexity – all without breaking the blind.



"I had never seen this capability before. We refer to it here as a 'non-interim interim,' and it has made a tremendous difference in our study analyses."

Chief Medical Officer, West Coast Biotech

PREVAIL UNIFIED ANALYTICS SELECTED CUSTOMER RESULTS

Leveraging insights from Prevail's innovative technology, customers have:

Achieved Fast Track designation and approval for a rolling NDA submission from the FDA, accelerating development timelines by several months

Had Phase II studies accepted as pivotal studies, saving millions in development costs and further reducing project timelines

Reduced patient ineligibility to less than 1%, with the study completed ahead of schedule and reaching all primary and secondary endpoints

Discovered promising new therapeutic indications in study data, with high potential commercial value

Avoided a clinical hold, saving $9 million in additional study costs

"We were up and running with integrated clinical, logistics, and financial oversight in 17 days for our large Phase III study, and enrolled the first patient on the 18th day."

Head of Clinical Operations,
East Coast Biotech

KEY STATS

Unify visualizations and reporting from

>15 Data Sources

Implement in less than

4 Weeks

PREVAIL ANALYTICS

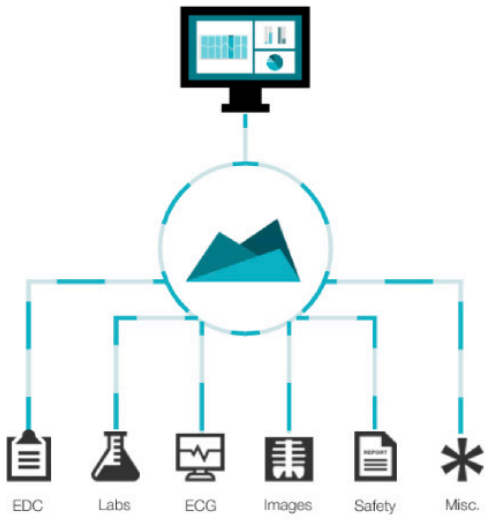
Integrated Analytics

Improve study success with early observations of aggregated data across studies and programs

Prevail's innovative analytics provide timely, data-driven insights into live studies, regardless of protocol, data, and system complexity – all without breaking the blind. With advanced visibility into aggregated clinical and operational data at their fingertips, sponsors are making faster and more-informed decisions to achieve their study endpoints, enhance patient safety, and reduce risk.

Deployed as Software-as-a-Service, Prevail Unified Analytics can be implemented in just 4-6 weeks, even in studies already underway – and is often up and running in just 15 days. All clinical, operational, diagnostic, and accounting data is automatically and continually integrated into a central repository for rapid and comprehensive analysis – from any source or system.

Clinical researchers thus get rapid, direct access to analytics and visualization tools without requiring IT or programming involvement, applying their knowledge and expertise to adjust data parameters on the fly and see the results immediately.



Source and system-agnostic, Prevail's technology easily aggregates and analyzes data from any system and source, including EDC, CTMS, labs, imaging, ECG, safety, IRT, ePRO, accounting, and more.

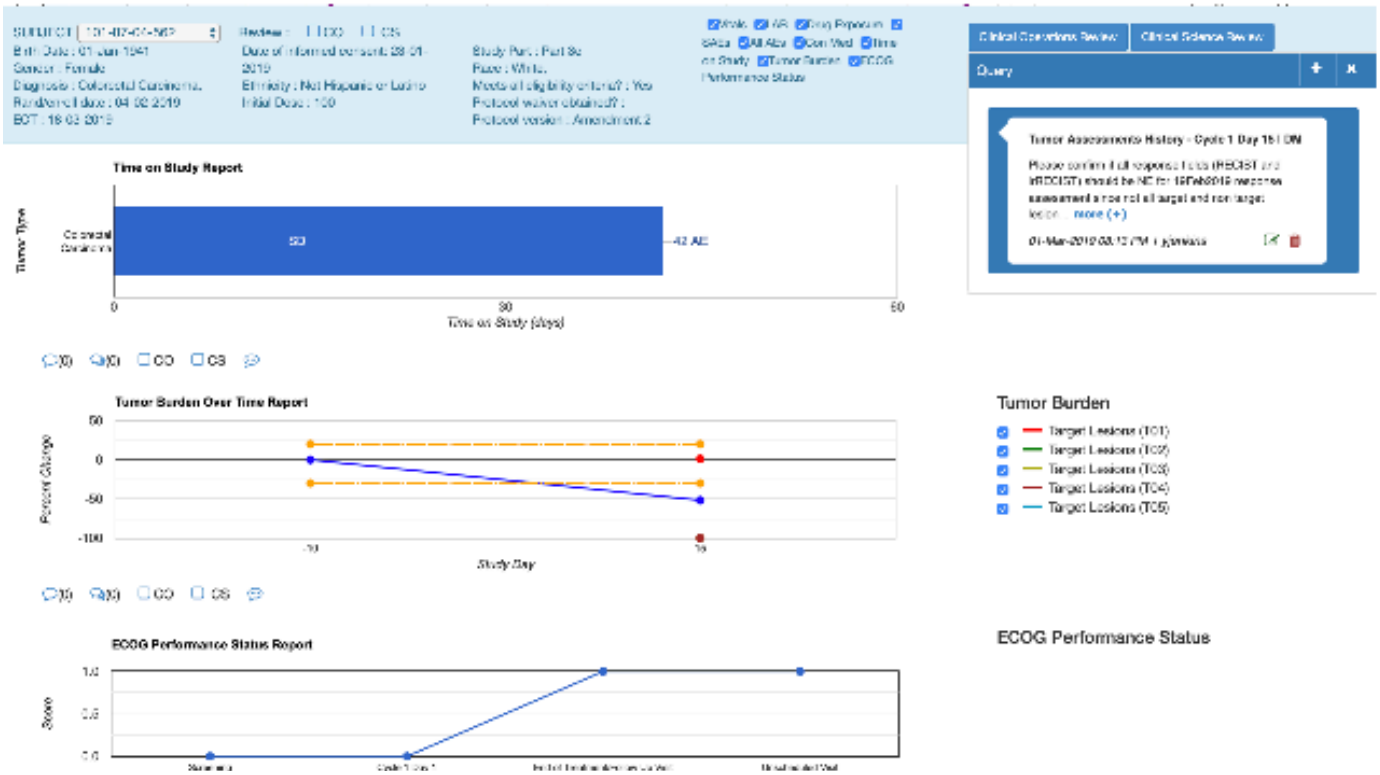
Prevail Unified Analytics: Key Features & Benefits

Accelerate Timelines

Quickly obtain data-driven insights into hidden trends, correlations, and outliers in aggregated data and workflow processes to enable agile clinical development, while streamlining operations and data flow across sites, systems, CROs, labs, and more. Easily provide the data and analysis necessary to support expedited regulatory programs such as Fast Track and rolling submissions.

Improve Outcomes

Obtain real-time, actionable intelligence into every aspect of your study - from operational metrics tracking, to visual patient profiles, to safety and efficacy trends, to data quality and compliance. Knowledge is always at your fingertips with comprehensive analytics and reporting, without requiring programmers or IT support.



Prevail's unique ability to provide a unified view of a study encompassing all clinical and operational data sources has helped customers reduce development timelines by months, and study costs by millions of dollars.

Reduce Cost

Leverage higher-quality data and analytics to reduce the need for additional trials. Increase internal efficiency and quickly implement protocol amendments with greater accuracy by integrating clinical data with workflows and processes. Maximize the benefits of your existing systems while reducing legacy overhead.

Minimize Risk

Get advance visibility across your clinical programs to identify problems sooner. Quickly verify study assumptions against real-time data to enhance patient safety, prevent quality problems, and avoid cost overruns and trial delays.

Fast Time-to-Value

Thanks to more than 100 pre-built connectors to commonly used vendor systems, you can start seeing the benefits of Prevail technology in as little as two weeks for simpler studies, and in six weeks or less for even the most complex protocols. Our innovative SaaS technology fully automates the data extraction process from any existing internal system, as well as external ones provided by CROs and other service providers - independent of whether a system supports APIs or SFTPs, and in any format (e.g. not only SAS).

PREVAIL INFOWORKS

RISE ABOVE

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About Prevail InfoWorks

Prevail is a pioneering life sciences software company with a unique combination of clinical expertise and engineering prowess, delivered through a best-in-class, modular eClinical ecosystem with sophisticated analytics and visualizations and supported by Prevail domain experts.

Our innovative and patented technology quickly and easily integrates, normalizes, reconciles, and presents aggregated data, analysis, trends, and metrics encompassing all study-related data sources through a single interface - making clinical development faster and easier, while reducing trial risk. With Prevail, trial sponsors can obtain unmatched, real-time answers to virtually any clinical, operational, and financial question regarding a study or program.