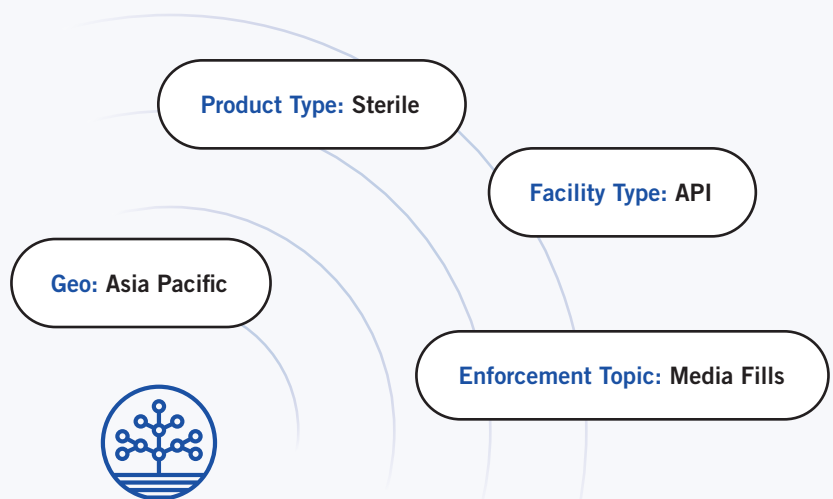

Redica Insights for Clinical Quality

How Can Redica Help You with Clinical Quality?

For firms involved with clinical development, how can you ensure quality? What if there was a tool to help you evaluate clinical sites, CROs, and IRBs? What if this tool could also provide real-time updates on the latest global regulatory developments affecting clinical research?

SEE THE WHOLE GCP PICTURE WITH OUR ACTIONABLE DATA INTELLIGENCE

We'll help you search, analyze, and derive valuable insight from thousands of CFR citations, 483s, and warning letters covering all GCP, GLP, and GVP areas.



THE REDICA DIFFERENCE

At Redica, we create actionable data intelligence that empowers global quality and compliance teams. Our Enforcement Analytics (EA) module addresses your deepest concerns around vendor quality, providing you the peace of mind to move your business relationships forward. Our database includes inspection history for hundreds of CRO, inspector profiles, and much more!

Redica's External Monitoring module provides automated, customized surveillance, expert analysis and machine translations under a single pane of glass. External Monitoring combines modern technology with the "real world" knowledge of seasoned experts who have defined this space for some of the largest companies in the world.

Learn more about how Redica Systems can help you!

A RESOURCE FOR CLINICAL QUALITY

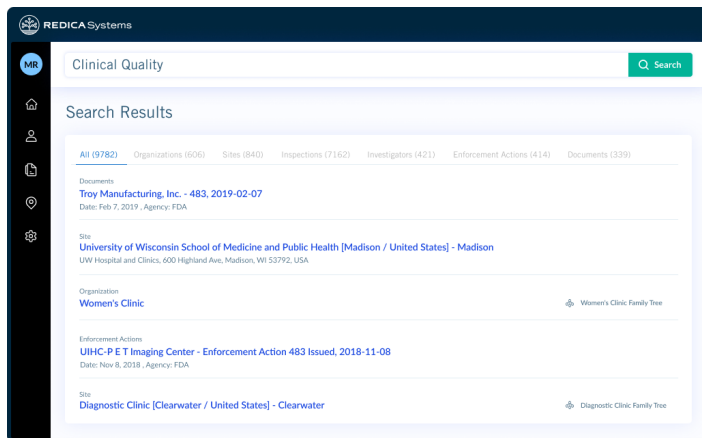
- Data on all FDA inspected and registered sites since 2000, with more than 350,000 sites and 800,000 inspections.
- Inspection reason for inspections—including preapproval, routine, and for-cause inspections.
- Over 100,000 tagged sites and organizations categories including sterile, API, FDF, CROs, clinical investigators, vaccines, etc.
- Expert tagging models for human drug GCP and medical devices that align citations with your quality system documents and SOPs.
- Updates on the latest regulatory developments plus analysis from leading industry experts.

REDICA INSIGHTS FOR CLINICAL QUALITY

Tailor insights to your specific compliance and risk concerns

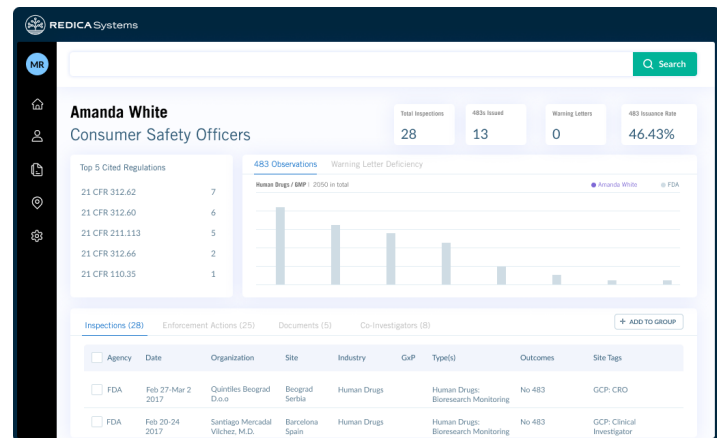
UNIFIED SEARCH

We'll help you search, analyze, and derive valuable insights from thousands of global regulatory data points covering CRO facilities, products, organizations, and enforcement.



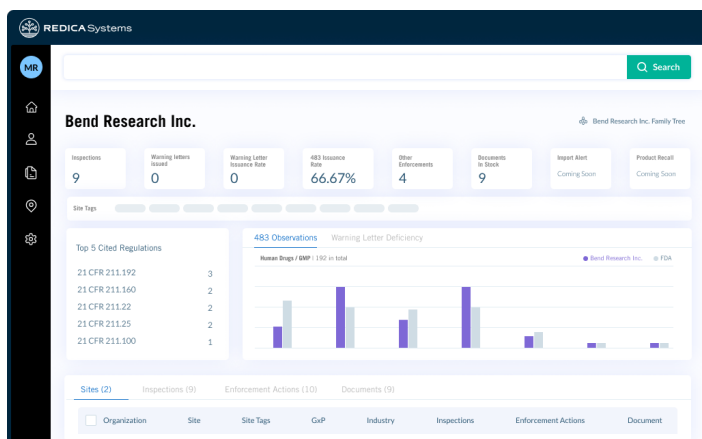
CLINICAL INVESTIGATOR PROFILES

The latest intelligence on health authority inspectors—their 483 observations, Warning Letter citations, co-inspectors, and more.



CRO ORGANIZATIONAL PROFILES

Our Family Tree allows you to see all sites, including suppliers and outsourcing facilities, associated with a particular organization in addition to inspection history, 483 observations, and Warning Letter citations.



SITE PROFILE

Inspection data for thousands of global sites, including inspection history, 483 observations, and Warning Letters. Plus, our Family Tree allows you to see all sites, including suppliers and outsourcing facilities, associated with a firm.

