REDICA | Monitoring

How do you stay on top of evolving regulatory changes to strengthen compliance?

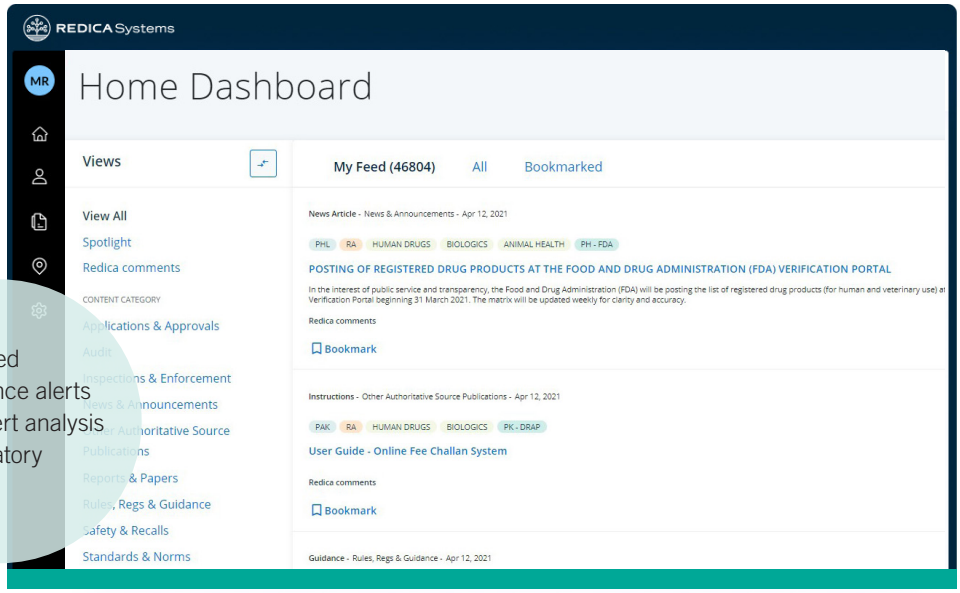
Redica's External Monitoring module provides automated and customized surveillance of changes in the regulatory environment, expert commentary, and machine translations to ensure signals aren't missed. Leveraging Redica's modern technology and the "real world" knowledge of seasoned experts gives you back precious time for analysis, planning, and taking critical action.

>65 Global Health Agencies

3,000 Data Sources

Expert Daily Monitoring

Streamlined Communication



Automated surveillance alerts and expert analysis on regulatory changes

STAY ABREAST OF CRITICAL REGULATORY SIGNALS

Traditional regulatory intelligence approaches are resource-intensive and still often miss risk signals and emerging trends. Streamline your regulatory intelligence process to ensure you never miss an important signal to remain compliant.

OUR CURRENT SOLUTION IS HIGHLY CONFIGURABLE

We configure our tools to your unique monitoring objectives and data requirements. Our dynamic design capabilities ensure each user can quickly find, review, and assess critical information based on their functional area.

WEEKLY SCANS AND HIGH IMPACT ALERTS FROM INDUSTRY EXPERTS

Monitoring communication includes links for drill-down research and commentary from Redica experts. Additional analysis opportunities are available for users with our Redica Systems Enforcement Analytics module.

REDICA Systems

OUR APPROACH

INTELLIGENCE GATHERED BY LEADING INDUSTRY EXPERTS

A critical differentiator in our External Monitoring solution is the combination of Redica's industry leading technology and automation with oversight and direct access to some of the best minds in the industry.

JANE WASTL

Senior GMP Quality Expert

Served in Eli Lilly leadership and technical positions in site and corporate quality assurance, quality control laboratory, training, master scheduling, capacity planning, and six sigma.

JAMIE COLGIN

GCP Expert

Experience in statistics, validation, audits, monitoring, validated SAS program development, GLP & GCP system evaluation, P&P writing, and set new standards for communicating audit findings.

BARBARA W. UNGER

Senior GMP Quality Expert Consultant

Led Amgen's Corporate GMP Audit group focused on API manufacturers, Quality Systems, and Computers. Developed, implemented, and maintained the GMP Regulatory Intelligence program for 8 years.

JERRY CHAPMAN

Senior GMP Quality Expert

Designed and implemented Eli Lilly's comprehensive GMP intelligence process to identify, analyze, and archive pertinent drug GMP regulations, inspection findings, trend discovery, and best practices.

MARK AGOSTINO

GMP/Medical Device/Combo Products Expert

Experience in quality assurance, supplier quality, and regulatory affairs.