Seeq® FOR PHARMA | ANALYTICS & AI SUITE

Advanced Data Analytics for the Life Sciences

Advanced data analytics in GMP Operations

From advancing active ingredients process development to maximizing fill-finish lines' throughput to streamlining regulatory reporting, Seeq empowers your subject matter experts with the data and capabilities they need to make better, faster decisions - the decisions that fuel innovation, ensure consistent quality and maximize production capacity.

The Seeq for Pharma Analytics & AI Suite is purpose-built to unleash the potential of under-leveraged time series data and to meet the unique requirements of the pharmaceutical industry. The suite is designed to be implemented in GMP manufacturing environments, providing robust analytics capabilities and features that support 21 CFR part 11 compliance and facilitate validation processes.

The biggest and most innovative Pharmaceuticals companies in the world use Seeq

6 of the top 10 global pharmaceuticals companies

30% of the top global contract manufacturers

Lilly · Abbott · Pfizer · Bristol Myers Squibb · Roche · Johnson & Johnson · Lonza

A platform for real-time data connectivity, analytics, transformation, and 21 CFR Part 11 compliance.

Data Integrity **Security** **Tracking**

Seeq FOR PHARMA ANALYTICS & AI SUITE

Your Data + Your Team = Your Advantage

Seeq empowers Pharma teams with intuitive, no-code analytics and GenAI, accelerating insights and improving quality while supporting 21 CFR Part 11 compliance through robust data integrity, audit trails and version control.

Seeq INDUSTRIAL ENTERPRISE MONITORING™

Scaled, Sustainable, System-Wide

Seeq simplifies Pharma monitoring with integrated insights, anomaly detection, and collaboration — enabling faster decisions, improved outcomes, and enhanced visibility across GMP processes, batch monitoring, and regulatory reporting.

Seeq AI Assistants speed adoption, skill building, & ROI

Driving Compliance and Efficiency

Seeq offers key features that support Pharma teams in ensuring data integrity, collaboration, and streamlined processes

Analysis Locking
Ensures production analysis remains unchanged

Analysis Versioning
Tracks and ensures transparency of analysis history

Readable Audit Trail
Provides rigorous documentation for all changes

Seeq SaaS Update Coordination
Supports software version updates for Pharma testing and validation

Seeq Additional Tenet
Dedicated test server for software validation

Right to Audit
Resources for quality audits and periodic reviews

Access to Pharma Trust Center
Portal for documentation, regulatory notes, and testing

Seeq Pharma Community of Practice
Peer networking, exclusive events, and participate in user research