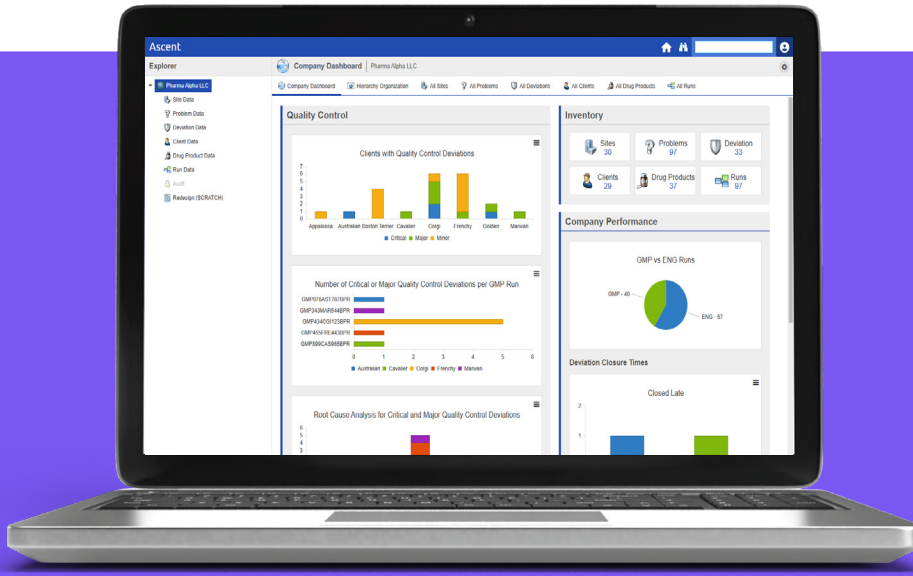
ASCENT

Addressing Challenges in Pharmaceuticals



Growing complexity within outsourced and distributed pharmaceutical manufacturing has revealed the limitations of legacy systems. Many contract development and manufacturing organizations (CDMOs) rely on cumbersome spreadsheets, disconnected tools and generic software, which are not optimized for manufacturing and industry good practices (GxP).

Client management and deviations often involve multiple applications and systems. This fragmentation causes friction for the business and critical workflows like batch record reporting.

For example, a client sampling data request may require many logins to find the proper data, searching through messages to management, a paper logbook and a shared drive. These delays can lead to client mistrust, slipping timelines and delayed decisions.

The Visiv platform addresses these critical challenges for pharmaceutical manufacturing. Visiv Ascent complements existing systems, centralizing GxP production run data and enabling deviation tracking and problem-solving based on real-time data from the manufacturing floor.

With our simple, purpose-built tool, CDMOs can onboard quickly, align on data and processes and empower employees to run faster with fewer surprises. Visiv Ascent lets users quickly and proactively address critical issues on the floor.

BENEFITS

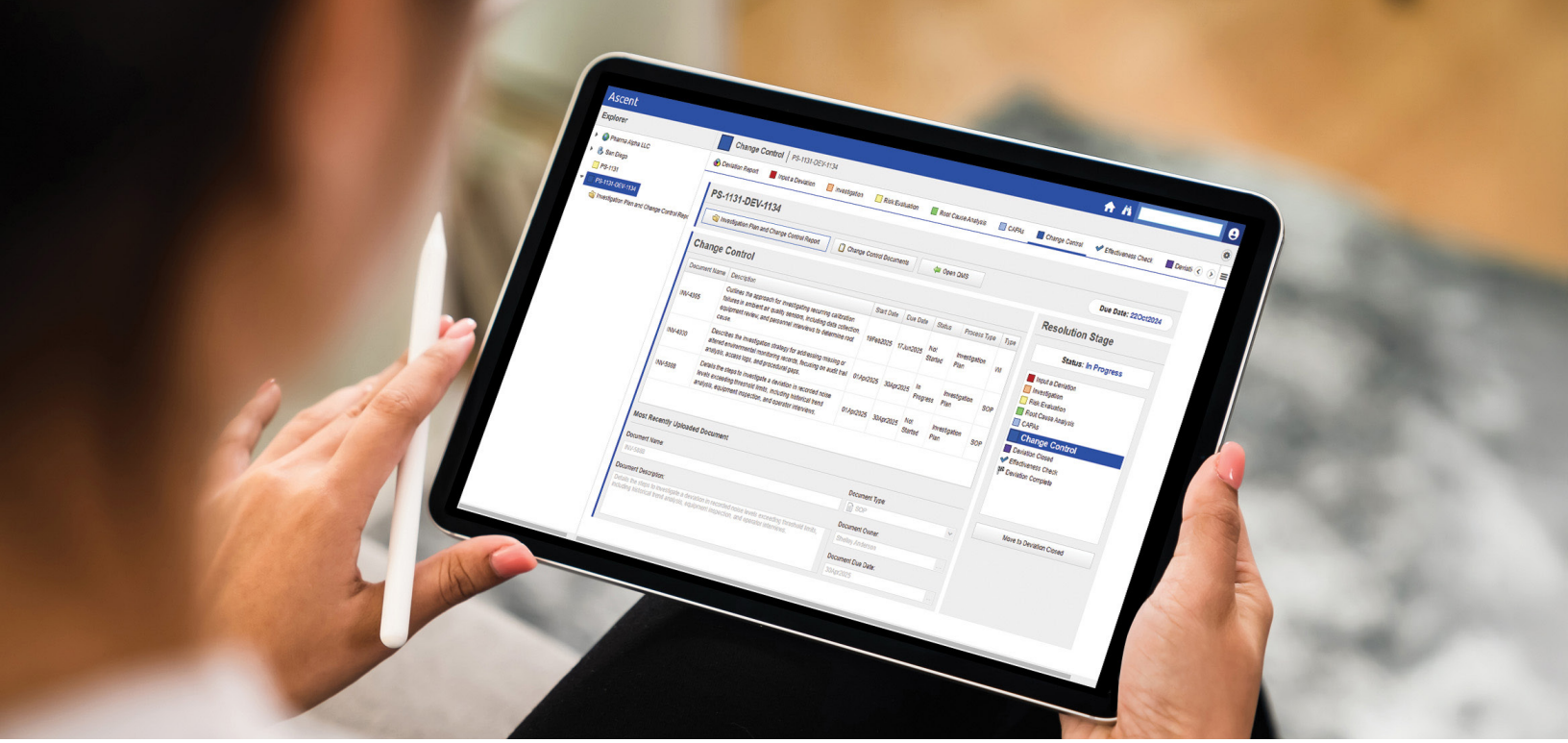
BATCH COORDINATION
Enable your teams—from production to quality—to make confident, real-time decisions that improve margins and client satisfaction.

BUILT FOR CDMOs
Deploy, configure and maintain quickly and easily—without sacrificing functionality, compliance or flexibility.

CLIENT AND AUDIT READINESS
Quickly answer client questions, support regulatory audits and uncover insights to optimize manufacturing timelines.

SCALABLE GxP PLATFORM
Adapt nimbly to changing client demands or regulatory shifts with a flexible, GxP-aware platform that supports rapid deployment and scale-up.

VISIV



FEATURES

AUDIT TRAILS
Track every data change, user action and process system event for full transparency and accountability.

CHART FILTERING
Drill into data by drug product, deviation, problem, site, production run or client, with no need for IT requests or complex workflows.

DASHBOARDS
Gain actionable insights into quality events on the manufacturing floor through intuitive visual dashboards.

DEVIATION MANAGEMENT
Monitor deviations across multiple drug products or clients. Spot trends by drug product, site or client to reduce repeat issues.

DOCUMENT MANAGEMENT
Identify and access documents needed to complete current Change Control and Investigation steps with built-in tracking.

MANUFACTURING PROCESSES
Define customized manufacturing process steps to fit your batch records.

REPORTS
Follow every step of the process with robust audit trails and reports, allowing visibility into the system anytime.

SAMPLING INFO
Enjoy the convenience of sampling logs purpose-built for export to clients when requested, with just one click.

VISIV'S PLATFORM

The power behind Visiv is a proprietary low-code / no-code platform and off-the-shelf industry applications that enable users to manage and leverage complex scientific or technical information to make confident, profitable business decisions, across portfolios, at any scale.

ABOUT VISIV

Visiv was founded to empower companies in highly scientific and technical industries to realize the full potential of their intellectual property, manage projects more efficiently and maximize investment value.

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