

TRANSFORM YOUR QC LAB

ACCELERATE BATCH RELEASE WITH BIOVIA

Optimize QC Labs • Ensure compliance • Maximize productivity

Challenges

Today's QC labs face pressure to quickly analyze the performance and quality of products, releasing batches as fast and cost-efficiently as possible. Traditional disconnected, paper-based, and error-prone processes cause:

Lengthy Delays

Up to
4 weeks
to release a
manufacturing batch

Up to
40 hours
to complete **batch review** processes

Compliance Issues

49%
of FDA warning
letters cite **data integrity issues**

25%
of lab expenses
spent **finding and correcting errors**

Increased Costs

45%
of R&D materials
are ultimately
wasted

Cost of **poor quality** up to
40%
of operating
expenses

The Solution

How to Accelerate Batch Release



**IF IT CAN BE
STANDARDIZED
IT SHOULD BE
STANDARDIZED**

Standards-based Methods
accelerate tech transfer and
time-to-market

Standardized procedures and data
ensure reproducible results and data re-use

**IF IT CAN BE
AUTOMATED
IT SHOULD BE
AUTOMATED**



Automated Review-by-Exception
to approve test results that meet
pre-determined tolerances

Automated Release-by-Exception
to instantly release for batches within
specifications

Value from BIOVIA

Productivity



25% reduction
in QA hours

60% reduction in
review times

95% faster data acquisition
from instruments

Quality & Compliance



Up to **80%** reduction in human errors

85% improvement in data
traceability

Costs



30% reduction in raw
material use

10% reduction in cost per
sample

BIOVIA ONE Lab

Up to **30%**
faster batch release
with Automated
Release-by-Exception

Up to **40%**
increase in efficiency with
guided procedure execution

Complete Integration:
of instruments and
materials in a unified
solution

Eliminate all paper:
including logbooks for
end-to-end traceability

LEARN MORE

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