BUTTERFLY

Chemicals

When the auditors arrive, are your records in one place?

In chemical manufacturing, when quality records, documents and approvals are spread across different systems, audit preparation turns into a project. Butterfly brings these processes into a single flow alongside your existing systems.

On-premises
deployment option

Without touching the ERP
alongside existing systems

Version history
on every document



Migrated with history

quality records arrive from the old system with their trail

PROBLEM

The challenges facing your organization

01 Deviations, change control and internal audit findings are tracked **in different systems and spreadsheets.**

02 Approvals wait **in email chains**; finding who approved takes time.

03 Before an audit, **record and document versions are gathered by hand.**

SOLUTION

How Butterfly solves it

Butterfly brings deviation, corrective and preventive action, change control and document processes together in a single orchestration layer. Every step is recorded with role, date and user; quality records in legacy systems can be migrated together with their history.

- ✓ Deviation, CAPA and change control in one flow
- ✓ Approvals securely between internal and external networks
- ✓ Maintenance work order created automatically on threshold
- ✓ Contract and document approval in the same layer

OUTCOME

What changes with Butterfly?

Audit ready
every step recorded with role and date

No code
rules are updated through data

Unbroken trail
from record to approval

This page is not a sector-specific customer case; it describes the capabilities of the Butterfly platform in this sector.

Do you have a similar process?

Let's look at it together in a 30-minute meeting.

Scan the QR code to request a meeting.

Request a meeting