PHARMACEUTICAL QMS SOFTWARE

Quality Management System (QMS)

Deviations, CAPA, change control, audits, and risk in a closed loop — every quality event connected, investigated, and closed.

OVERVIEW

GAMP QMS keeps every quality event in one system, linked to the events around it. Open a record and you can see what caused it, what it caused, and who signed each step and when.

WHY TEAMS CHOOSE GAMP QMS

Close the loop on every event

A deviation is raised, investigated and closed in one place, and the CAPA it triggers is attached to it — including the check that the action actually worked.

Control what changes

Change control with impact assessment, and audits handled in the same system, so a change and the audit that examines it are never separate records.

See risk before it becomes a finding

Risk assessment, out-of-specification and out-of-trend results, complaints and vendor oversight in one view of where quality is exposed.

CAPABILITIES

- ✓ Deviation handling with root cause analysis
- ✓ CAPA with effectiveness checks
- ✓ Change control and impact assessment
- ✓ Internal and external audit management
- ✓ Risk assessment, OOS/OOT, market complaints
- ✓ Vendor qualification and oversight

ON THE COMPLIANCEONE PLATFORM

- Single sign-on across modules, managed through one administration layer
- Active Directory integration
- Audit trail, electronic signatures and role-based access
- Cloud, on-premises or hybrid deployment
- Built for 21 CFR Part 11 and EU Annex 11

GAMP Lifetech Private Limited
sales@gamplifetech.com · +91 90160 72807
www.gamplifetech.com/platform/complianceone/quality-management-system

Issued November 2026