bizzmine QHSE

CAPA MANAGEMENT

CAPA management enables you to continuously improve the effectiveness of your quality management system.

ENHANCED COMPLIANCE

- **Comprehensive problem management**: Document non-conformances and quality alerts. Capture all relevant data.
- **In-depth analysis**: Analyse problems identify root causes.
- **Risk assessment**: Evaluate the potential impact of identified issues and prioritise actions.
- **Continuous improvement**: Measure and validate the effectiveness of implemented CAPAs.
- **Customisable audit forms**: Tailored audit forms to specific needs.

INCREASED FOLLOW-UP

- **Effective CAPA tracking**: Identify, analyse, and track deviations from audits and other compliance checks
- **Loopback mechanism**: Revisit the issue and refine the approach if a CAPA is found to be ineffective.
- **Audit Trail**: Maintain a complete audit trail of all CAPA activities, documenting every step.
- **Consistent review**: The systematic follow-up process reinforces accountability and ensures that corrective actions lead to lasting improvements.

REDUCED MANUAL WORK

- **Automated CAPA processes**: Focus on higher-level tasks, knowing that the system will handle routine activities reliably.
- **Task reminders**: Assigned users are automatically reminded of tasks they need to complete on time.
- **Streamlined initiation of CAPA**: Let you guide through the necessary steps, across different teams and departments.
- **Efficient audit management**: Automated tracking of audit findings with direct link to corrective actions.

IMPROVED DECISION-MAKING

- **Informed action planning**: Make more informed decisions when planning and implementing corrective actions.
- **Customisable dashboards**: Tailored dashboards to display KPIs, audit results, or the status of ongoing CAPAs.
- **Continuous Monitoring**: Go for improvement through real-time feedback on the success of implemented actions.
- **Root Cause Analysis**: Generate detailed data on root causes using predefined categories.

> *With the Bizzmine workflow module, the CAPA and MoC processes have been made paperless and are compliant with the requirements of 21 CFR Part 11.*
> *- **Carbogen AMCIS***

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CORRECT DEVIATIONS, ENABLE CONTINUOUS IMPROVEMENT

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