

## MPS 6 – Quality Assurance and Process Integrity

**\*\*Category:\*\*** Process Integrity

**\*\*Tags:\*\*** process integrity, quality assurance, procurement controls, behavioral compliance, change management, anomaly detection, loss prevention, sabotage risk, continuous improvement

**\*\*Description:\*\*** Minimum Performance Standard for Quality Assurance and Process Integrity. Defines the intent, required actions, and guidance for establishing reliable, auditable processes across all operational domains — including procurement, HR, manufacturing, logistics, and finance. This MPS promotes loss prevention, anomaly detection, and continuous improvement through structured process design, control gap analysis, behavioral compliance, and change management.

### Assessment Criteria (Structured)

1. 1.

**\*\*Requirement:\*\*** Key business processes must be mapped and documented, including inputs, outputs, decision points, and control owners.

**\*\*Evidence:\*\*** Approved process maps for core functions such as procurement, onboarding, payroll, manufacturing, and logistics.

2. 2.

**\*\*Requirement:\*\*** Control gaps must be identified and classified using historical losses, audits, feedback, and testing.

**\*\*Evidence:\*\*** Documented gap analyses and control failure classifications.

3. 3.

**\*\*Requirement:\*\*** Quality assurance mechanisms must be implemented to verify compliance with intended process flows.

**\*\*Evidence:\*\*** Checklists, approval logs, exception reports, and spot-check records.

4. 4.

**\*\*Requirement:\*\*** Behavior-based compliance monitoring must be used to detect informal workarounds and SOP deviations.

**\*\*Evidence:\*\*** Observational compliance reports, incident logs, and escalation records.

5. 5.

**\*\*Requirement:\*\*** Formal change management protocols must govern updates to systems, processes, and procedures.

**\*\*Evidence:\*\*** Change request forms, approval workflows, and user communication logs.

6. 6.

**\*\*Requirement:\*\*** Anomaly detection systems must monitor for process bypasses, manipulation, and output deviations.

**\*\*Evidence:\*\*** System alerts, anomaly reports, and monitoring dashboards.

7. 7.

**\*\*Requirement:\*\*** Preventative and loss mitigation measures must be documented and monitored.

**\*\*Evidence:\*\*** Trigger thresholds, early warning indicators, and access control reviews.

8. 8.

**\*\*Requirement:\*\*** Independent internal QA audits or reviews must be conducted periodically.

**\*\*Evidence:\*\*** QA audit reports, corrective action plans, and adherence assessments.

9. 9.

**\*\*Requirement:\*\*** Line managers and process owners must be trained to detect control failures and drive remediation.

**\*\*Evidence:\*\*** Training logs, session materials, and feedback forms.

10. 10.

**\*\*Requirement:\*\*** A central repository must store all process maps and QA documentation in an accessible format.

**\*\*Evidence:\*\*** Centralized library with indexed process maps, SOPs, and QA standards.