SOP: Follow-up Inspections and Quality Assurance Checks

This SOP details procedures for **follow-up inspections and quality assurance checks**, ensuring consistent compliance with established standards, identifying and addressing deficiencies, maintaining product and service quality, documenting findings, and implementing corrective actions to enhance overall operational effectiveness and customer satisfaction.

1. Purpose

To outline standardized procedures for conducting follow-up inspections and quality assurance (QA) checks, ensuring sustained compliance and continual improvement.

2. Scope

This SOP applies to all departments/units responsible for product or service delivery requiring ongoing quality oversight.

3. Responsibilities

- QA Team: Conduct inspections and QA checks, document findings, and recommend corrective actions.
- Department Managers: Ensure implementation of corrective actions and ongoing compliance.
- Staff: Cooperate during inspections and participate in corrective measures as required.

4. Procedure

1. Initial Inspection Review

Review prior inspection/QA reports to identify areas requiring follow-up.

2. Scheduling

Determine schedule for follow-up inspections or QA checks based on risk, recurrence of issues, or regulatory requirements.

3. Preparation

Prepare inspection checklists, prior findings, and necessary materials.

4. Conducting Inspection/QA Check

- Perform physical inspections, process audits, or service reviews as applicable.
- Use standardized checklists for consistency.
- Engage relevant staff as needed.

5. Documentation

- Record findings using standardized forms or digital systems.
- Document deficiencies and non-compliances in detail.

6. Reporting

- Prepare and distribute inspection/QA reports to relevant stakeholders.
- Highlight unresolved or recurring issues.

7. Corrective Actions

- $\circ\;$ Assign responsibilities and deadlines for corrective actions.
- o Track progress and verify completion in subsequent follow-ups.

8. Continuous Improvement

- Analyze trends in findings and corrective actions.
- Update processes and training as necessary.

5. Documentation and Records

Document/Record	Location	Retention Period
Inspection/QA Checklists	QA Shared Drive	3 years
Inspection/QA Reports	QA Shared Drive	3 years
Corrective Action Logs	QA System	3 years after closure

6. Review and Revision

- This SOP shall be reviewed annually or upon significant process change.
- Revisions must be documented and dated in the change log.

7. References

- Company Quality Policy
- Regulatory and industry standards as applicable

8. Change Log

Date	Version	Description	Author
2024-06-05	1.0	Initial release	QA Department