

SQP Document List

- 1. Organization chart
- 2. Responsibility and/or job description,
- 3. Quality System Procedures (e.g., quality policy, objectives, manual and procedures for the Quality Management System and other processes)
- 4. Management review records
- 5. Internal audit documents (audit plan, report, etc.)
- 6. Supplier Control documents (supplier approval procedure / criteria, list of approval supplier list, supplier evaluation records, on-going performance monitoring, etc.)
- 7. Document control procedure and records (including record keeping)
- 8. Product specifications/requirements
- 9. Inspection Instructions, acceptance criteria and inspection & testing reports (including the stages of IQC, In-process and Final inspection)
- 10. Work instructions / workmanship standards for each manufacturing process
- 11. Production schedules/records
- 12. Procedure for defining and reporting of "incident"
- 13. Product recall procedure
- 14. Customer complaints records
- 15. Corrective action reports (related to incident, internal audit, complaint, etc)
- 16. Test records on Traceability system
- 17. Equipment maintenance documents (plan, procedure, record, etc)
- 18. Calibration of monitoring & measuring devices (plan, procedures, records, etc)
- 19. Cleaning schedule and procedure
- 20. List of Approved Chemicals with Corresponding Brands / Manufacturers
- 21. Pest control documents (list of trained pest control staff, contract with external pest control agency, pest control inspection record, bait documentation, etc)
- 22. Record / plan for "Risk Assessment" of the entire manufacturing processes
- 23. Risk assessment records of final product
- 24. Product testing procedure/program
- 25. Laboratory test reports (including lead and heavy metals content in paints, coatings and non-paint components, hardware, labels, final product, etc).
- 26. Monitoring records of foreign body detectors (e.g. metal detection records, daily sensitivity checking records of metal detectors...etc)
- 27. Broken needle procedure & records (if applicable)
- 28. Pre-production meetings records
- 29. Process Control Plan
- 30. Training (procedure, training needs & records)

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Page 1 of 1

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