Manufacturing Quality Nonconformance Investigation Report

Report Number:

Date:

Prepared By:

Department:

Nonconformance Description

Product/Part No.:

Batch/Lot No.:

Description:

Detected By:

Detection Date:

Investigation Details

Root Cause Analysis:

Immediate Actions Taken:

Disposition and Corrective Actions

Disposition: Select

Corrective/Preventive Actions:

Responsible Person(s):

Completion Date:

Verification of Effectiveness

Verified By:

Verification Date:

Verification Outcome:

Signatures

Name	Signature	Date