

# Corrective and Preventive Action (CAPA) Plan Template

<b>CAPA Number</b>	[Enter CAPA ID]
<b>Date Opened</b>	[Enter date]
<b>Responsible Person</b>	[Name/Title]
<b>Department</b>	[Enter department]

## 1. Description of Issue / Nonconformity

[Provide a concise description of the problem, nonconformity, or incident. Include relevant references or identifiers as needed.]

## 2. Root Cause Analysis

[Summarize analysis performed to determine the root cause(s) of the issue. Methods such as 5 Whys, Fishbone Diagram, etc. can be referenced here.]

## 3. Corrective Action Plan

### Action(s) to Correct Immediate Issue

[List steps taken or to be taken to resolve the current issue.]

Action Step	Responsible Person	Due Date	Status
[Step 1]	[Name/Role]	[Date]	[Open/In Progress/Done]
[Step 2]	[Name/Role]	[Date]	[Open/In Progress/Done]

## 4. Preventive Action Plan

### Measures to Prevent Recurrence

[List process changes, training, audits, controls, or other long-term prevention measures that will be implemented.]

Preventive Action	Responsible Person	Due Date	Status
[Measure 1]	[Name/Role]	[Date]	[Open/In Progress/Done]
[Measure 2]	[Name/Role]	[Date]	[Open/In Progress/Done]

## 5. Effectiveness Verification

[Outline how and when you will verify that corrective and preventive actions were effective. Specify criteria, methods, responsibilities, and timelines.]

## 6. Closure

<b>Reviewed By</b>	[Name/Title]
<b>Date Closed</b>	[Enter date]
<b>Comments</b>	[Any additional notes or comments]

## **Important Notes**

- CAPA plans are critical for managing quality and continuous improvement.
- All actions and investigations should be evidence-based and well documented.
- Responsibility and timelines must be clearly assigned and tracked.
- Verifying effectiveness ensures that similar issues do not recur.
- Maintain CAPA records as part of your compliance and audit readiness.